

#### **ASX Release**

#### **Suda Presentation**

Stephen Carter, Executive Chairman for Suda Ltd, is currently meeting with brokers, carrying out a road show in Perth, Sydney and Melbourne.

A copy of the presentation follows.

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# SUDALTD

Stephen Carter Executive Chairman /CEO

# **SUDALTD**

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SUDA Ltd (ASX:SUD) is an Australian Drug **Delivery company focused on oro-mucosal** delivery of existing drugs using our novel delivery technology (SUDAMist<sup>™</sup>). The acquisition of NovaMist<sup>™</sup> technology complements this model and provides a pipeline of products in various stages of development. This acquisition assists SUDA in its goal to become a major player in oromucosal products.

### ArTiMist™

- Our lead product ArTiMist™, a sub-lingual artemether based malaria treatment, has been developed specifically for children with complicated malaria.
- In 2011 Thompson Reuters identified ArTiMist™ as one of the worlds 5 most promising drugs in Phase III clinical development.

 Successfully completed ArTiMist<sup>™</sup> Phase III
clinical trial in Children in Severe or complicated Malaria.



### ArTiMist™

#### Phase III Results

- 151 patients in 3 sites in Africa (Rwanda, Ghana and Burkina Faso)
- Sub-lingual ArTiMist<sup>™</sup> superior to i.v. quinine
- 95.6% of ArTiMist<sup>™</sup> patients had reduced parasite counts by >90% in the first 24 hours compared with only 40.6% using intravenous (i.v.) quinine
- Parasite clearance time 30 hours with ArTiMist<sup>™</sup> vs. 68 hours with i.v. Quinine
- No Early treatment failures in the Artimist<sup>™</sup> arm 10 in quinine arm.
- Results support that ArTiMist<sup>™</sup> is an superior treatment for children with malaria
- Results provide a compelling argument for the potential use of ArTiMist<sup>™</sup> as an early interventional treatment for children

### ArTiMist™

Advantages of ArTiMist<sup>™</sup> over other treatments are:

- It does not require hospitalisation or medically trained personnel for administration;
- Not affected by GI complications;
- By-passes the Liver and the significant metabolism seen from the first pass effect;
- Does not require fatty diet for maximum effect;
- Rapidly absorbed;
- Can be administered to comatose patients;
- Negates risk of infection from needle use;
- Has a long shelf life; and
- Critically, in hot climates, does not require cold chain storage

### ArTiMist™ Next Steps

- Final Report due July 2013
- Discussions with potential trade sale partners
- Finalise the Regulatory documents Dec 2013
- Filing of Regulatory documents Jan 2014
- Trade sale

#### **ArTiMist**<sup>™</sup> Value Proposition

- Phase III trails confirm superiority of ArTiMist<sup>™</sup> over gold standard i.v. quinine.
- Successful commercial scale manufacture program completed.
- Drafting of Regulatory documentation advanced.
- Turn key trade sale of asset proposed.
- Massive un-met medical need.

#### NovaMist™

- SUDA has signed a Sale and Purchase Agreement to purchase the NovaMist<sup>™</sup> IP and inventory.
- The agreement is subject to Novadel shareholder approval and should be finalised Mid July 2013.
- The benefit of this technology is that it not only compliments SUDA's current ArTiMist<sup>™</sup> Project but it also provides a strong foundation for the development of other drug candidates which may open the doors to lucrative licensing agreements.

#### NovaMist™

The NovaMist<sup>™</sup> platform provides a valuable life cycle extension strategy for innovators whose products are facing patent expiration and new product development opportunities for companies wishing to develop their own brand of a competitor's drug that is approaching patent expiration.



The key product in development, Duromist<sup>™</sup>, is a stable solution of lingual Sildenafil (the active ingredient in Viagra<sup>™</sup>) that has shown preliminary bioequivalence to Viagra<sup>™</sup> tablets in early clinical trials. The Erectile Dysfunction market worldwide is \$4.1 billion and outside of the USA is approx. \$2billion pa

#### **NovaMist**<sup>™</sup>

- NovaDel's NovaMist<sup>™</sup> technology may provide substantial potential benefits compared to other modes of drug administration including:
  - Faster onset of action
  - Lower dose
  - Enhanced patient compliance and convenience
  - Avoiding the need to swallow
  - Allowing medication to be taken without water
  - Increased bioavailability of drug by avoiding metabolism by liver

#### NovaMist™

- There are a number of products in various stages of development including.
- Note all the following products have shown good results in proof of concept studies.
  - sumatriptan, (worlds largest selling migraine drug) being developed for the treatment of migraine headache); (Phase 2/3)
  - Duromist<sup>™</sup> (an oral spray formulation of sildenafil, being developed for the treatment of erectile dysfunction); (Phase 1/2)
  - Zensana<sup>™</sup> (an oral spray formulation of odansetron, being developed for the treatment of chemotherapy-induced nausea and vomiting); (Phase1/2)
  - NVD-101 (oral spray formulation of sildenafil being developed for the treatment of Pulmonary Arterial Hypertension) (Phase 1/2)
  - NVD-301 (an oral spray formulation of midazolam, being developed for sedation during diagnostic, therapeutic, and endoscopic procedures). (Pre-Clinical)

### SUDALTD Product Pipeline



SUDA is progressing rapidly to commercialisation of the worlds first sublingual Malaria treatment for children.

### **SUDALTD** Future milestones

Milestone	Expected time frame
ArTiMist™ Final Clinical trial Report	July 2013
Completion of NovaMist™ Acquisition	July 2013
odging of ArTiMist™ Regulatory File in first jurisdiction.	December 2013 - January 2014
First NovaMist™ deal	March -April 2014
ArTiMist™ Trade Sale	February-August 2014

## SUDALTD Our People

#### Stephen Carter. Chairman and Chief Executive Officer

 Has extensive pharmaceutical industry experience with multi-national pharmaceutical and listed public companies.

#### Michael Stewart. Non-Executive Director

 Broad corporate and management background and involvement in bilateral donor funded and World Bank co-financed Aid Projects

#### Ken Robson Non-Executive Director

 His background includes extensive experience as a Corporate Lawyer and Advisor, specialising in fundraising, market compliance and Mergers & Acquisitions.

#### Joseph Ohayon CFO and Director

 Has over 20 years' experience in financial roles including 12 years within health-related industries.

### SUDALTD Corporate Information

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ASX Code:		(ASX:SUD
Market Capitalisation:		\$20 M
Number of Shares on Issue:		626,091,2
6 month high:	30/04/2013	\$0.049
6 month low:	16/11/2012	\$0.027
30 Day VWAP:		\$0.037
Average Trading Volume:		1.7M
Cash at 1 <sup>st</sup> April 2013		\$1.3M

# SUDALTD Thank You