

Extraordinary General Meeting

Sophia Antipolis – October 13, 2015

This presentation contains certain forward-looking statements. Although the Company believes its expectations are based on reasonable assumptions, these forward-looking statements are subject to numerous risks and uncertainties, which could cause actual results to differ materially from those anticipated in the forward-looking statements.

Risk factors which are likely to have a material effect on Nicox's business are presented in the 4th chapter of the "Document de référence, rapport financier annuel et rapport de gestion 2014" filed with the French *Autorité des Marchés Financiers* (AMF) on April 10, 2015, which is available on Nicox's website (www.nicox.com).



Building a leading Global Ophthalmic Company



BAUSCH + LOMB



1 Vesneo™: potential to become the leading intraocular pressure (IOP)-lowering drug in the glaucoma space

- Partnered worldwide with Bausch + Lomb (Valeant)
- NDA filed with FDA – PDUFA date **July 21, 2016**
- Peak sales guidance of >\$500 million in U.S. and >\$1 billion WW¹
- Potential net milestones up to \$132.5 million and 6%-11% net royalties
- Engine for NO-donating pipeline growth

Product	Region	Phase	Regulatory Status	Commercial Status	Notes
AC-170	US	Phase 1	Completed	Completed	Patented cetirizine eye drop formulation for allergic conjunctivitis; potential FDA approval expected by end 2016.
NCX 4251	US	Phase 1	Completed	Completed	Patented nanocrystalline fluticasone for blepharitis; expected to enter US phase 2 trials post-IND approval.
Next generation NO-donors	US	Phase 1	In Progress	In Progress	Targeting glaucoma and reduction of IOP

2 Further advanced proprietary pipeline

- **AC-170**: patented cetirizine eye drop formulation for allergic conjunctivitis; potential FDA approval expected by end 2016.
- **NCX 4251**: patented nanocrystalline fluticasone for blepharitis; expected to enter US phase 2 trials post-IND approval.
- **Next generation NO-donors** targeting glaucoma and reduction of IOP

3 European commercial business

- Track record of revenue growth from diversified product portfolio
- Product launches planned throughout 2015-2018
- AzaSite® and BromSite™ MAA filing planned by H1 2016



1. Valeant corporate press release 09/26/14

Vesneo™: Significant revenue potential for Nicox

- U.S. glaucoma market valued at nearly \$2.4 billion annually¹
- Partnered with Bausch + Lomb, a division of Valeant Pharmaceuticals
 - Provided **peak sales guidance of >\$500 million in U.S., >\$1 billion WW²**
 - Expands Bausch + Lomb's presence in chronic eye diseases
- Potential revenues from worldwide licensing agreement with Bausch + Lomb
 - \$20 million paid to Nicox between 2010 and 2012 (upfront + 1st milestone)
 - Remaining milestones up to \$162.5 million (\$132.5 million net)
 - \$30 million due to Pfizer in two payments, with majority of commercial milestones to Nicox
 - Potential tiered royalties to Nicox from 6% - 11% (net of single digit royalty to Pfizer)

NDA filed by Bausch + Lomb – PDUFA date July 21, 2016

1. IMS December 2014 (USC #61690 oph prostaglandins and miotics, #61660 oph beta blockers, #61650 oph carbonic anhydrase inhibitors)
2. Valeant corporate press release 09/26/14

Vesneo™: Clinical Program

- Discovered and synthesized in Nicox Research Labs in Milano
- First NO-donating compound for IOP lowering in patients with open angle glaucoma or ocular hypertension
- Robust clinical program
 - Phase 3 Studies
 - Completed two large international pivotal Phase 3 efficacy studies (APOLLO and LUNAR) with 840 patients overall in U.S. and Europe
 - Studies met their primary endpoint and showed positive results on a number of secondary endpoints
 - Phase 2 Studies
 - Completed four Phase 2 studies
 - Statistically significant greater IOP lowering versus Xalatan® (latanoprost) in a Phase 2b dose ranging study (VOYAGER)
 - Significant improvement in 24-hour IOP lowering versus timolol in sleep lab Phase 2 study (CONSTELLATION)

No significant safety issues noted as of top-line results Phase 2 or Phase 3

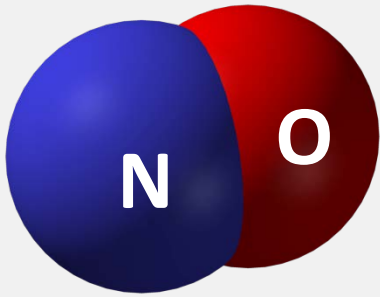
Robust unencumbered pipeline beyond VESNEO

Product	Rights	Preclinical	Development	Regulatory/Marketing	Status
Core worldwide pipeline					
AC-170 (cetirizine) <i>Ocular itching associated with allergic conjunctivitis</i>	Worldwide	→			Potential FDA approval by end 2016
NCX 4251 (fluticasone propionate nanocrystals) <i>Blepharitis</i>	Worldwide	→			Expected to enter phase 2 pending IND approval
NCX 470 (NO-bimatoprost) <i>Glaucoma</i>	Worldwide	→			Preclinical
Next generation NO-donors <i>Glaucoma and other ophthalmic indications</i>	Worldwide	→			Lead optimization
European pipeline					
AzaSite® (1% azythromycin) <i>Bacterial conjunctivitis</i>	EMEA ¹	→			European filing expected by H1'16
BromSite™ (0.075% bromfenac) <i>Pain and inflammation after cataract surgery</i>	EMEA ¹	→			European filing Expected by H1'16
NCX 4240² (Carragelose) <i>Viral conjunctivitis</i>	Worldwide	→			European launch expected in 2017
AAT³ (RPS-AP) <i>Diagnostic test for the combined detection of adenoviral and allergic conjunctivitis</i>	Worldwide	→			

1. Europe Middle East Africa 2. Medical device 3. In vitro diagnostic medical device



Our proprietary Nitric Oxide donating platform



Nobel Prize 1998
RF Furchgott
LJ Ignarro
F Murad

Endogenous cell-signaling molecule

Nitric Oxide (NO) in ophthalmology

NO and other molecules involved in NO-mediated signaling are present in ocular tissues. NO notably plays a role in the regulation of intraocular pressure (IOP). This is of particular interest in glaucoma, which is often associated with increased IOP.

- **First generation NO-donors**

- Designed to release both NO and a well-established drug
- Potential to provide enhanced efficacy compared to the well-established drug

VESNEO™
latanoprostene bunod

NCX 470

- **Second generation stand-alone NO-donors**

- Enable optimization of NO dosing
- Target ophthalmic diseases which have a major component modulated by NO

NCX 667

5-for-1 reverse stock split

All shareholders will automatically receive 1 new Nicox share in exchange for 5 existing Nicox shares.

- If approved by the shareholders and implemented by the Board of Directors, the reverse stock split is expected to become effective in approximately 2 months.
- A reverse stock split will reduce the number of outstanding common shares from 114.3 million to 22.9 million.
- A detailed calendar will be communicated in due time.

What will I have to do?

- Shareholders who hold a total number of shares that is an exact multiple of 5 will not need to take any action, and the reverse stock split will be carried out automatically.
- Shareholders who do not hold a total number of shares that is an exact multiple of 5 will have to buy or sell existing shares to reach a multiple of 5. Alternatively, remaining fractional shares will be sold by their financial intermediary and shareholders will receive a cash payment.

Completion of a reverse stock split is a critical step in a larger strategic initiative to increase awareness and visibility for Nicox ahead of the potential FDA approval of VESNEO next year.



Key upcoming milestones

Clinical & Regulatory

- ❑ Vesneo™ (latanoprostene bunod) – PDUFA date July 21, 2016
- ❑ AC-170 – Potential FDA approval expected by end 2016

Corporate

- ❑ Continued execution of EU commercial plan with positive sales momentum
- ❑ Additional European product launches 2015-2018, including AzaSite® and BromSite™
- ❑ Further in-licensing and corporate development opportunities



Nicox

Visible science

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