



## 2009 Half year report – Telephone conference

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# Forward looking statements



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- **Half year report 2009**
- **Highlights**
  - New partner collaborations
  - Pipeline news
- **Pipeline status**
- **Outlook and expectations**



# Financial reporting H1 2009



(DKK million)	<b>H1 2009</b>	H1 2008	FY 2008
Revenue	<b>19</b>	33	67
Cost	<b>(223)</b>	(221)	(433)
Operating loss	<b>(204)</b>	(188)	(366)
Finance, net	<b>12</b>	(6)	(21)
Results affiliates	<b>(7)</b>	(6)	(29)
Tax	<b>21</b>	14	34
Net result	<b>(178)</b>	(186)	(382)
Capital resources	<b>489</b>	716	481

- Full year guidance unchanged: Operating loss ~ DKK 350 mill.
- Capital resources unchanged through first half of 2009
- Capital resources increased in August 2009 to ~ DKK 730 mill.

# Financing: Commercial transactions in 2009



- **GSK – January; expansion of former agreement**
  - Upfront payment + milestones and royalties
  
- **Eli Lilly – February; new drug discovery alliance**
  - USD 30 m in guaranteed funding + milestones and royalties
  
- **GSK – August; Advance of NSD-721 into Phase I**
  - Exercise of EUR 5 mill. share put option
  - Cash milestone payment of EUR 4 mill.
  
- **Janssen – August; new drug discovery alliance**
  - EUR 32 mill. in guaranteed funding + milestones and royalties

## Conclusions and outlook

- Four partner transactions in 2009
- Attractiveness of CNS Drug Discovery
- Financing ~ DKK 500 mill.
- Significant future revenue potential
- Strategic and financial goals obtained

# Main highlights in 2009



- **Huntexil™ (pridopidine); lead orphan drug for Huntington's disease**
  - Completed recruitment in MermaiHD, a European Phase III HD study (437 patients)
  - Launch of Compassionate Use programme in Europe
  - The HART study in NA is still recruiting patients, while progressing satisfactorily
  
- **Tesofensine; Best in class anti-obesity drug (NCE)**
  - Successful outcome of End of Phase II meeting with the FDA
  - Intensified partner discussions in parallel with final Phase III preparations
  
- **ACR343; Supportive results in Phase I. Prepared for Phase II in schizophrenia**
  
- **ACR325; Prepared for first patient study in Parkinsons Dyskinesia**
  
- **NSD-788; Positive Phase I results and Proof-of-mechanism in depression/anxiety**
  
- **NSD-721 (GSK); Successfully advanced into Phase I (EURm 9 financing from GSK)**
  
- **3 new partner deals: Guaranteed funding of ~DKK 500 million + significant potential**

# Pipeline



Indication	Programme	Mechanism of Action	Partner	PC dev.	Phase I	Phase II	Phase III	NDA / Reg.
Huntington's disease	Huntexil™	Dopaminergic stabil.						
Obesity	Tesofensine	MRI						
ADHD	ABT-894	NNR modulator	Abbott					
Dyskinesias (PD)	ACR325	Dopaminergic stabil.						
Schizophrenia	ACR343	Dopaminergic stabil.						
Cognitive dysfunctions	ABT-560	NNR modulator	Abbott					
Depression/anxiety	NSD-788	MRI						
Social anxiety	NSD-721	GABA modulator	GSK					
Schizophrenia	NSD-761	Ion channel mod.						
Autoimmune diseases	NSD-726	Ion channel mod.						
Psychoses	NSD-847	Dopaminergic stabil.	GSK					
ADHD	NSD-867	Cortical enhancer	GSK					

# Expectations next 6 months



- Huntington's disease - Huntexil™ (pridopidine)
  - First results from pivotal Phase III programme
- Obesity - Tesofensine
  - Finalise Phase III preparations
  - Partner discussions
- Dyskinesias in Parkinson's disease - ACR325
  - Initiation of Phase Ib study in Parkinson's patients with dyskinesias
- Schizophrenia - ACR343
  - Initiation of Phase II studies
- Further pipeline advances
- Additional partner agreements





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