

Photocure ASA

- from Biotech to Specialty Pharma

Presentation of fourth quarter 2009 results

19 February 2010

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Highlights fourth quarter 2009

- Sale of Metvix/ Aktilite for EUR 51 million 30 September 2009
- Received positive response to the Hexvix NDA from FDA on 30 December 2009
- Payment of dividend of NOK 4 per share in December 2009
- Cash of NOK 403.5 million as per 31 December 2009
- Sales revenues of NOK 14.6 million in Q409 (33.2)

Major steps towards a Specialty Pharma Company



From Biotech to Specialty Pharma

- We will create a unique Specialty Pharma Company
- We will continue to focus on PDT based products
- We increase our commitment to the Dermatology market
 - Capitalize on technological and commercial capabilities
- We remain committed to our Cancer portfolio
 - Continue providing PDT products for diagnosis & treatment

Technology and capabilities to create significant value



Strong platform for future growth

Creating value from strong IP position in Dermatology and Cancer

	Indication	Status	Peak sales potential EU/ US
Hexvix [®]	Detection of bladder cancer	Approved in EU Approval pending US	EUR 130 – 240 million
Cevira™	Treatment of cervical cancer	Phase I/II	EUR 250 – 550 million
Lumacan™	Detection of colon cancer	Phase I/II	EUR 300 – 510 million
Allumera™	Improvement of facial skin appearance	Pilot trial	EUR 30 – 50 million
Visonac™	Treatment of acne	Phase II	EUR 240 – 420 million

A promising pipeline with large market potential and well-defined roadmap to market



Financial statements

Profit & Loss

Q4 and Full Year 2009



- Net gain on sale of Metvix/Aktilite of MNOK 369.3 in 2009
- Reduced revenues and M&S expenses due to sale of Metvix/Aktilite
- R&D expenses of MNOK 27.3 in 4Q due to Hexvix NDA expenses and Visonac clinical phase II study

Numbers in NOK thousand	Q4 2009	Q4 2008	FY 2009	FY 2008
Total revenues	14 566	33 190	98 798	102 220
R&D expenses	-27 289	-25 508	-79 492	-78 341
Marketing & sales expenses	-8 012	-12 559	-41 640	-45 916
Operating profit/ loss (EBIT)	-29 918	-14 469	-59 394	-62 539
Net profit/ loss	-28 587	-9 246	-56 943	-59 562
Gain on sale of Metvix/ Aktilite	6 338	0	369 325	0
Net profit/loss	-22 249	-9 246	312 382	-59 562
Other comprehensive income	3 144	0	4 192	0
Comprehensive income	-19 105	-9 246	316 574	-59 562

Profit & Loss (cont.) Continued vs discontinued operations 2009



	Discontinued	l operations	Continued operations		
Numbers in NOK thousand	2009	2008	2009	2008	
Total revenues	50 370	65 366	48 428	36 854	
Gross profit	40 443	50 667	42 887	32 48	
Other income	0	0	11 652	3 580	
Indirect manufacturing expenses	-1 165	-1 694	-9 451	-6 913	
R&D expenses	-512	-1 816	-78 980	-76 52	
Marketing & sales expenses	-16 656	-21 642	-24 984	-24 27	
G&A expenses	-2 552	-362	-20 075	-16 040	
Operating profit/ loss (EBIT)	19 558	25 153	-78 953	-87 692	
Net profit/ loss	19 558	25 153	-76 501	-84 71	
Gain on sale of Metvix/ Aktilite	369 325		0	(
Discontinued operations	-388 883	-25 153	388 883	25 15	
Net profit/loss	0	0	312 382	-59 562	

Metvix/Aktilite is under IFRS reported as discontinued operations



Segment information – Q4 2009

(unaudited) Q4 2009				Q4 2009				800	
Numbers in NOK thousand	Own	Partner	R&D*	Total	% vs. 08	Own	Partner	R&D*	Total
Sales revenue Metvix/ Aktilite	-307	2 913		2 606	<i>-</i> 87 %	7 421	12 223		19 644
Sales revenue Hexvix	3 843	8 117		11 960	-12 %	3 368	10 178		13 546
Total sales revenues	3 536	11 030	0	14 566	-56 %	10 789	22 401	0	33 190
Cost of goods sold	-145	-1 961		-2 106	-64 %	-810	-5 104		-5 914
Gross profit	3 392	9 069	0	12 460	-54 %	9 980	17 297	0	27 276
Gross profit	96%	82%		86%		92%	77%		82%
Gain sale Metvix/Aktilite	0	6 338		6 338		0	0		C
Operating expenses	-6 734	-7 905	-27 739	-42 378	2 %	-10 729	-3 236	-27 780	-41 745
Operating profit	-3 342	7 502	-27 739	-23 579		-749	14 060	-27 780	-14 469
* Including share of general & administrative expenses									

Hexvix sales increased 8% in end user sales and -12% in sales revenue due to Nordic inventory variations and COGS/currency adjustment related to Partner sales



Segment information – FY 2009

(audited)	FY 2009						F	Y 2008			
Numbers in NOK thousand	Own	Partner	R&D*	Disc. op.	Total	% vs. 08	Own	Partner	R&D*	Disc. op.	Total
Sales revenue Metvix/ Aktilite	0	2 430		50 370	52 800	-18 %	0	0		64 063	64 063
Sales revenue Hexvix	16 908	29 090		0	45 998	25 %	10 190	26 664		0	36 855
Milestone revenue	0	0			0		0	0		1 303	1 303
Total revenues	16 908	31 520	0	50 370	98 798	-3 %	10 190	26 664	0	65 366	102 220
Cost of goods sold	-718	-4 823		-9 927	-15 469	-19 %	-558	-3 817		-14 699	-19 074
Gross profit	16 190	26 696	0	40 443	83 329	0 %	9 632	22 847	0	50 667	83 147
Gross profit (ex milestones)	96%	85%		80%	84%		95%	86%		78%	81%
Operating expenses	-21 193	-17 908	-82 737	-20 885	-142 723	-2 %	-15 683	-13 786	-90 703	-25 514	-145 686
Operating profit	-5 003	8 788	-82 737	19 558	-59 394		-6 051	9 062	-90 703	25 153	-62 539
Gain sale Metvix/Aktilite	0	0	0	369 325	369 325		0	0	0	0	0
Profit before tax	-5 003	8 788	-82 737	388 882	312 382		-6 051	9 062	-90 703	25 153	-59 562
* Including share of general & adr	* Including share of general & administrative expenses										

Hexvix sales increased 18% in end user sales and 25% in sales revenue in 2009

Lower level of R&D expenses and total operating expenses in 2009 vs. 2008



Balance sheet – assets

- NOK 403.5 million in cash year end
- No Metvix/Aktilite assets
- Other investments includes1,040,000 shares in PCI Biotech valued at NOK 11.00/share

Numbers in NOK thousand (audited)	31.12.2009	31.12. 2008
Non-current assets		
Intangible assets, software	365	534
Machinery & Equipment	1 772	3 939
Other investments	14 585	11 528
Total non-current assets	16 722	16 001
Current assets		
Inventory	13 826	12 792
Receivables	22 811	29 158
Cash & cash equivalents	403 502	179 897
Total current assets	440 140	221 846
Total assets	456 862	237 847

Balance sheet - equity & liabilities



- NOK 415.8 million in shareholder's equity or 91%
- No interest bearing debt
- No Metvix/Aktilite liabilities

Numbers in NOK thousand (audited)	31.12.2009	31.12.2008
Paid-in capital	11 047	11 047
Other paid-in capital	176 112	15 467
Retained earnings	228 624	173 181
Shareholders' equity	415 783	199 694
Total equity	415 783	199 694
Total long-term liabilities	340	0
Accounts payable	13 936	0
Tax and social charge	3 325	0
Current liabilities	23 478	38 153
Total liabilities	40 739	38 153
Total equity and liabilities	456 862	237 847

Cash Flow Q4 2009 and Full Year 2009



- NOK 263.2 million in Net change in cash during Q4
 - Gain from sale of Metvix/Aktilite
 - Payment of dividend
 - Purchase of own shares

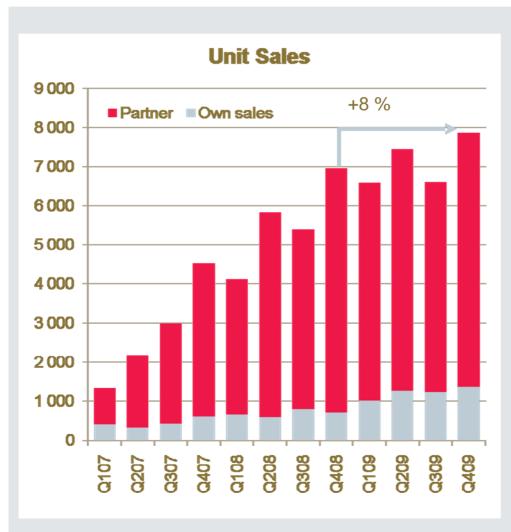
Numbers in NOK thousand	Q4 2009	FY 2009	FY 2008
Income/ loss before tax	-22 248	312 382	-59 562
Other operational items	385 022	5 063	3 584
Net cash flow from operations	362 774	317 445	-55 978
Cash flow from investments	4 357	10 104	-11 865
Payment of dividend	-87 950	-87 950	0
Purchase own shares	-19 915	-19 915	0
Sale of own shares	3 926	3 926	0
Interest paid	-1	-4	-13
Cash flow from capital transactions	-103 941	-103 944	-13
Net change in cash during the period	263 190	223 605	-67 856
Cash & cash equivalents beginning of period	140 312	179 897	247 753
Cash & cash equivalents beginning end period	403 502	403 502	179 897



Operational Update



Hexvix® key sales figures



Unit sales Q4 2009:

- 1,362 Hexvix units sold in the Nordic region, an increase of 30% vs. Q4 2008
- 6,503 Hexvix units sold by GE Healthcare, an increase of 4% vs. Q4 2008

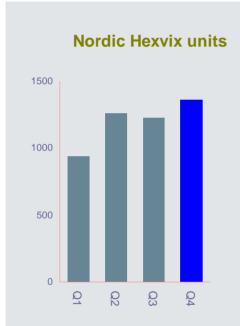
Unit sales 2009:

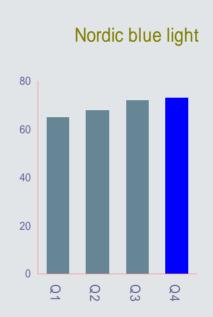
- 4,786 Hexvix units sold in the Nordic, an increase of 61% vs. 2008
- 22,962 Hexvix units sold by GE Healthcare, an increase of 18%vs. 2008

Hexvix®



- Nordic key performance indicators







Hexvix Nordic market share in TURB 23%

Equipment growth 18% in 2009

Growth in use 36% in 2009



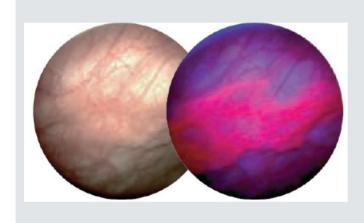
Hexvix® partner status 2009



- Germany largest market over 50% of sales
 - New reimbursement from 1. January 2010
- App. 600 blue-light scopes in market
- Established national and regional guidelines as well as EU consensus
- Focus on sales execution



Hexvix® roadmap



- Secure US approval
 - Recommended by FDA Advisory Committee and positive response, including action letter received from FDA in December 2009
 - Pending issues expected to be agreed during 1H 2010
- Launch in US in 2010
- Increase use in Europe
 - Several programs for training/reimbursement/sales execution/equipment placement depending on country

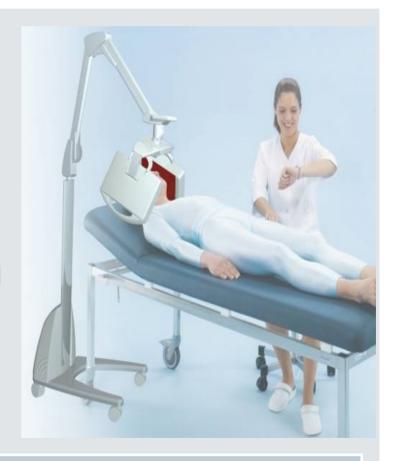
Peak sales potential of EUR 130 - 240 million in the EU & US

Visonac™



Effective treatment of moderate to severe acne

- 3 modes of action:
 - Kills the acne bacteria
 - Reduces sebaceous production
 - Reduces inflammation
- Consistently high efficacy shown in 3 separate Phase II studies
- Treatments two weeks apart sustained effect in reduction of lesions
- Limited side effects
 - Significant improvement in tolerability measures of pain and erythema



Peak sales potential of EUR 240 – 420 million in EU & US

Visonac[™]

Status and road map



- Completed patient enrollment in multicenter phase II study in US/Canada in January 2010
 - Pediatric patient population (n=107)
 - Preliminary results in March/April 2010
 - Study 2 months ahead of plan
- Results from phase II study will improve phase III program in EU/US
- Plan End-of-Phase II meeting with the FDA in the US and regulatory update in EU in Q3 2010
- Plan start of joint Phase III program in EU/US in Q4 2010

Alignment of EU and US program

AllumeraTM Improving facial skin appearance



- Cosmetic product sold through dermatologists
- Finished pilot study in Q4 2009 with excellent results;
 - Texture: Softened and smoothened skin
 - Tone: Evened out skin color
 - Fine lines: Diminished
 - Pores: Reduced pore lines
- Initiation of consumer trial in US in Q2 2010
 - Planned results for Q4 2010
- Started preparations for launch in 2011

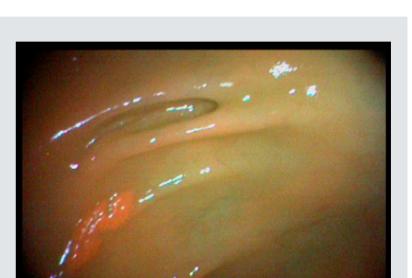


Peak sales potential estimated to be EUR 30-50 million per year in US

Lumacan™

Diagnosis of colorectal cancer

- Proof-of-Concept study using enema showed 39% increase in detection rate using Lumacan- colonoscopy.
- →Improved oral formulation in development to optimize release in colon
- → Scintigraphy testing of new oral formulations in Q2 2010
- Ongoing phase I/II study on hold after 12 patients in Q4 2009. Restarting with improved oral formulation in Q3 2010



First PoC-study in Munich, Germany. One flat lesion showing fluorescence in colon. Courtesy: Prof. Dr. B. Mayinger

Peak sales potential of EUR 300 – 510 million in the EU & US

CeviraTM



Treatment of HPV/precancerous lesions in cervix

- Completed Proof-of-Concept study in 2009 showed high efficacy in low grade pre-cancerous lesions
- Placebo-controlled multicenter phase II study ongoing in 5 countries in Europe
 - All 70 patients enrolled
 - Patient follow-up 6 months
 - Initial results expected Q2 2010
- Developed new drug/medical device
 - One visit to gynecologist
 - User friendly for patients disposable
- New phase II study in same population testing the device planned for H2 2010



Illustration of Cevira device

Peak sales potential of EUR 250 - 550 million in the EU & US



Summary



Goals 2010

Strategic:

Create a Specialty Pharma Company in dermatology – starting with the US

Commercial:

- Improve commercial activities for Hexvix in Europe
- Secure Hexvix Approval in the US
- Hexvix launch in the US

R&D:

- Start Visonac phase III program
- Finish Allumera consumer trial
- Cevira start phase II study with new device
- Lumacan restart phase II study with improved oral formulation