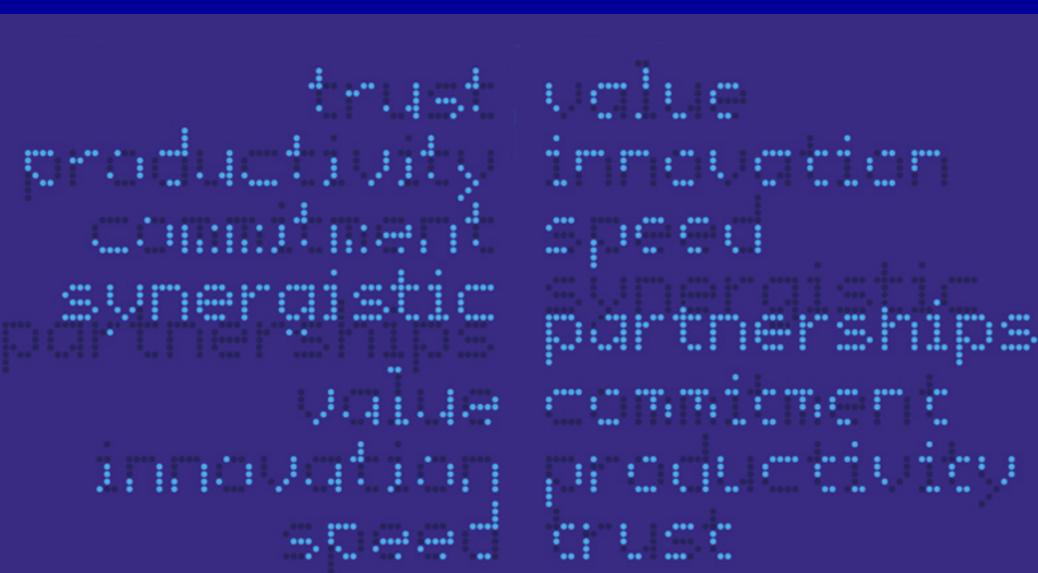


Evotec OAI AG Analyst Meeting, 26 March 2004





We attribute our successful 2003 to reliable and long-term partnerships





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- 01 Overview
- 02 Commercial Review
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Defying market trends in 2003

We delivered:

- Group revenue growth of 10% to EUR77.2m
- Positive Group-EBITDA
- Positive operating income before amortisation in Discovery and Development Services
- Tools and Technologies break-even
- Numerous new or extended customer programmes:
 Roche, Novartis, Oxford Bioscience Partners portfolio companies, etc.
- Implementation of Discovery Programs strategy:
 - Initiated strategic alliance with DeveloGen (Metabolic Diseases)
 - Signed € 20m contract with Takeda (Alzheimer's Disease)
 - In-licensed 5 pre-clinical compounds from Roche (CNS)



We are bridging the gap between early discovery and clinical trials for our partners





Why do our partners need us?

Biotech

Disease biology

- extensive genetically derived disease knowledge
- sub-critical discovery process know how
- lack of chemical libraries

Need to partner



Need to partner

- growing focus on Phase II/III + marketing
- increasing early stage in-licensing
- concentration on fewer suppliers

Development Marketing Pharma



Ideally suited to bring what our partners need

Biotech

Disease biology

- extensive genetically derived disease knowledge
- sub-critical discovery process know how
- lack of chemical libraries

Need to partner

Evotec OAI



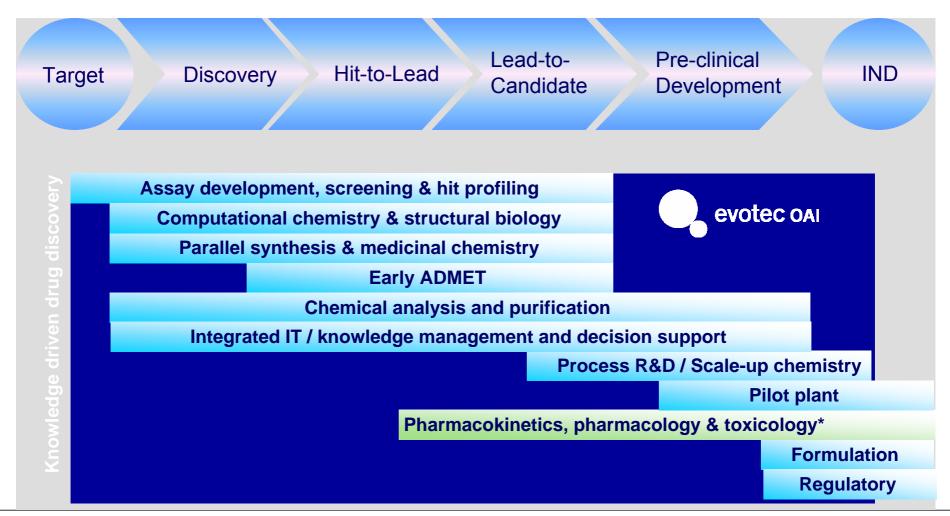
Need to partner

- growing focus on Phase II/III + marketing
- increasing early stage in-licensing
- concentration on fewer suppliers

Development Marketing Pharma



Critical success factor 1: Fully-integrated R&D solutions



^{*} typically done by customer or subcontracted



Critical success factor 2: Critical mass in science skills and facilities



- 27 biology and 5 cell-culture labs;>35 chemistry labs; 8 analytical labs
 - World-class screening factory:
 3 uHTS platforms for biochemical and cellular assays
 - 2 cGMP kilo laboratories and 2 pilot plants (FDA and European quality standards)
- Cutting-edge analytical equipment
- World-class knowledge management system
- 500+ experienced scientific staff:
 - > 300 discovery, development and analytical chemists
- Strong emphasis on speed, quality and customer service with a proven ability to execute



Critical success factor 3: Outstanding customer references

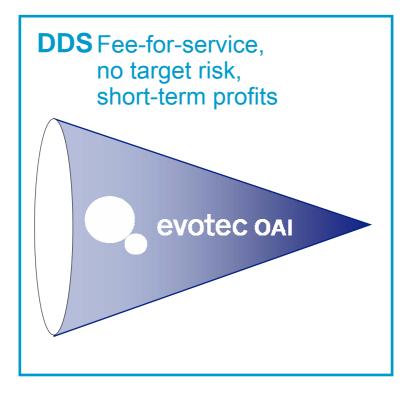


So far, we have delivered to our clients:

- > 20 lead compounds
- > 10 pre-clinical development candidates
- 3 clinically approved compounds



We offer fee-for-service projects (DDS) and run internal discovery programmes (DPD)

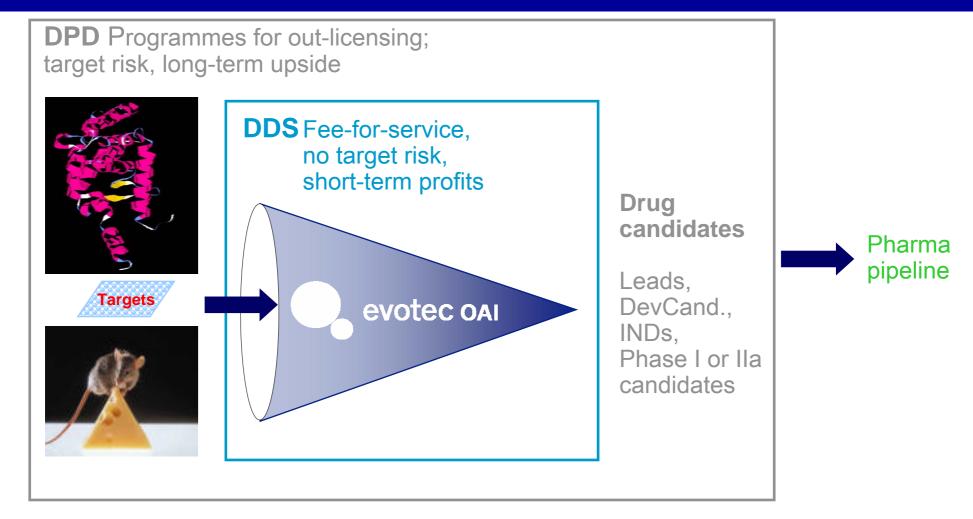


Assays, Screens, Libraries, Med Chem, PRD, Scale-up, Formulation





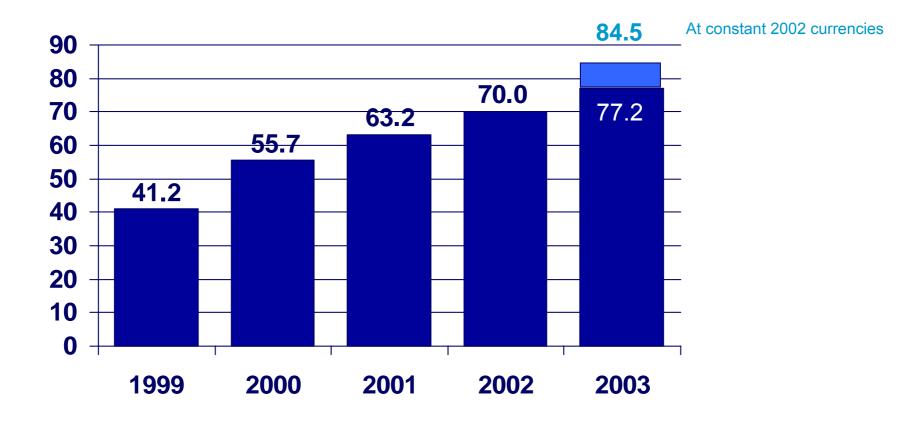
We offer fee-for-service projects (DDS) and run internal discovery programmes (DPD)





17% proforma growth over past five years In constant €s: 21% growth 2003 / 2002

EUR million





Financial highlights 2003: Strong growth, EBITDA profitability and stable cash position

	2003	2003
		Constant 2002 currencies
Revenues	EUR 77.2m	EUR 84.5m
Growth over 2002	10%	21%
- DDS	EUR 61.2m	EUR 68.5m
- DDS over 2002	4%	17%
Gross margin	40%	42%
EBITDA	EUR 4.1m	EUR 6.6m
Cash as of 31/12	EUR 19.5m	



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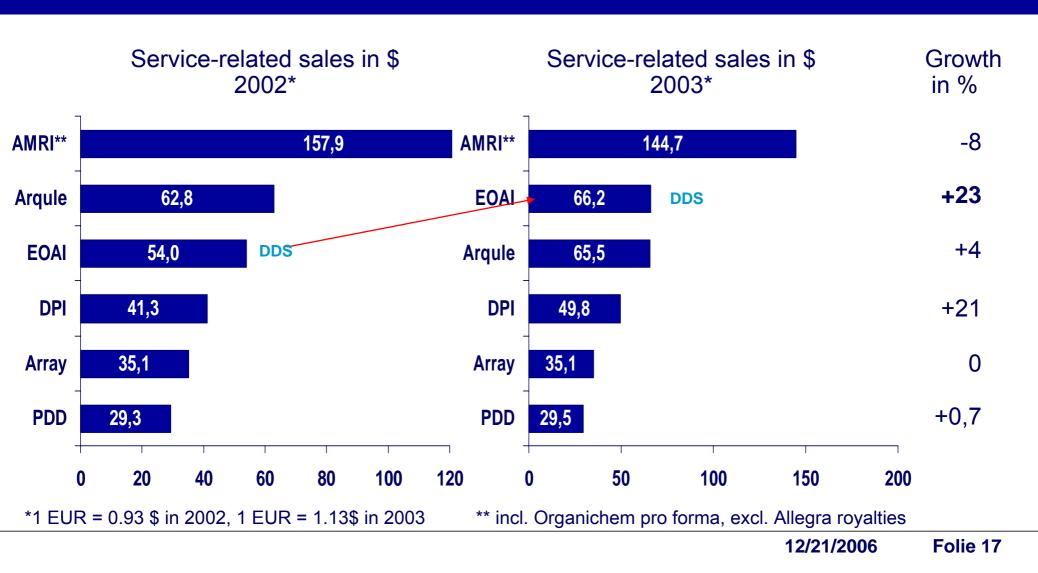


Commercial highlights

- Market share gain vs. main competitors in "Discovery and Development Services"
- Market share gains in "Tools and Technologies"
- Leading deal flow: within DDS > 30 new customers
- Long-standing customer base including most top pharma and biotech companies
- Repeat business growing by >10% (expansions of Roche, Novartis and >20 FTEs in OBP-related deals)



Market share gains in a tight market 14% in 2002 to 17% in 2003 ...

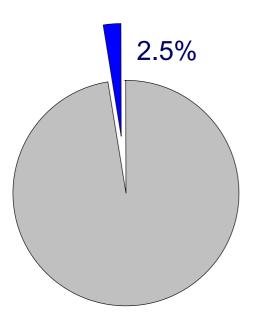


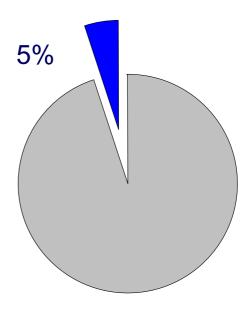


... and Evotec Technologies doubled their market share

ET sales 2002: \$10.6m







Total market* size ~ 400m US\$

Source: HighTech Business Decisions, Dec. 2003 *Total Market = High Throughput Screening for Detection Instruments, Liquid Handling and Automation Systems



A full range of integrated drug discovery and development solutions ...

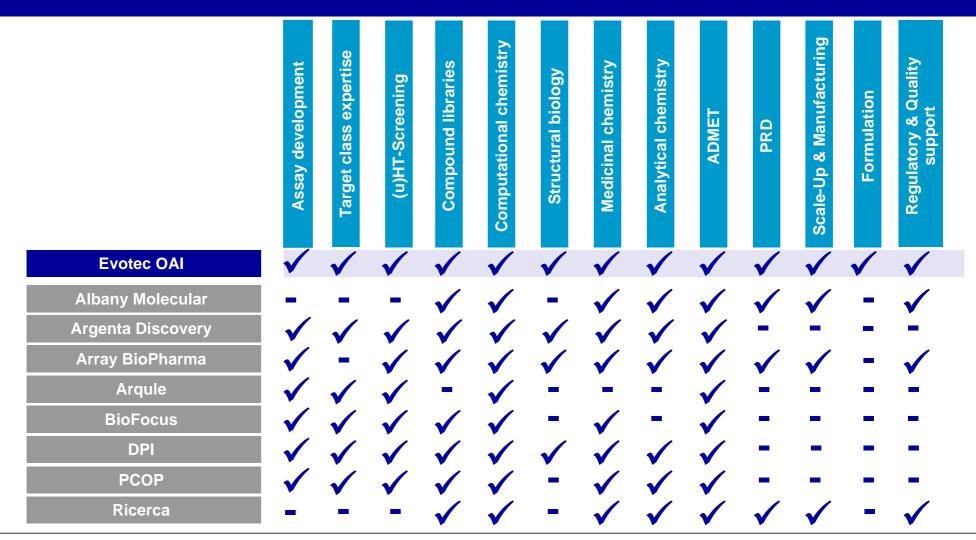


Highlights 2003

- GPCR collaboration with Euroscreen
- Expansion of cellular assay capabilities
- Build-up of Structural Biology & Computational Chemistry
- Expansion of ADMET platform
- Natural product collaboration with Biofrontera
- Full integration of ProPharma formulation capabilities



... providing a significant competitive advantage



Source: Company websites



We complement our customers' disease expertise





A long-standing customer base including most top pharma and biotech companies



































































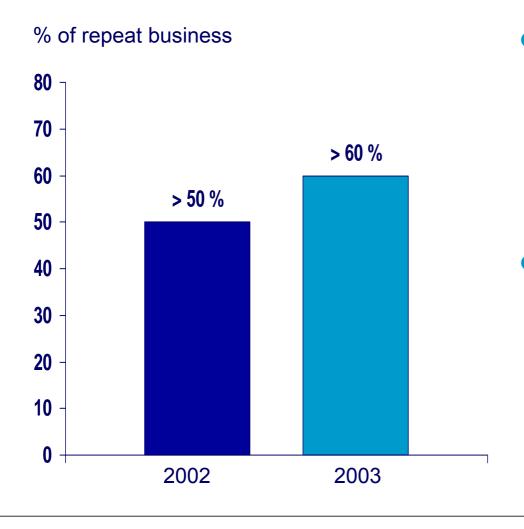


Major achievements (1)

- > 250 projects completed
- Long-term relationships
 - Grew collaborations with emerging biotech companies such as Biolmage, Elixir, Infinity, Oxigene and Prolysis in the areas of assay development, screening and medicinal chemistry
 - Strengthened relationships with Alizyme, Celgene, DuPont and Serono in development chemistry
- New customers
 - Secured > 30 new customers in 2003
 - Initiated collaborations with several pharmaceutical and biotechnology companies using our full range of integrated capabilities, e.g. Dynogen, Oxagen and Toray
 - Started new collaborations with Anormed and Stiefel in development chemistry
 - Build CNS partnership with Takeda



Major achievements (2) Novartis & Roche agreements expanded



Novartis

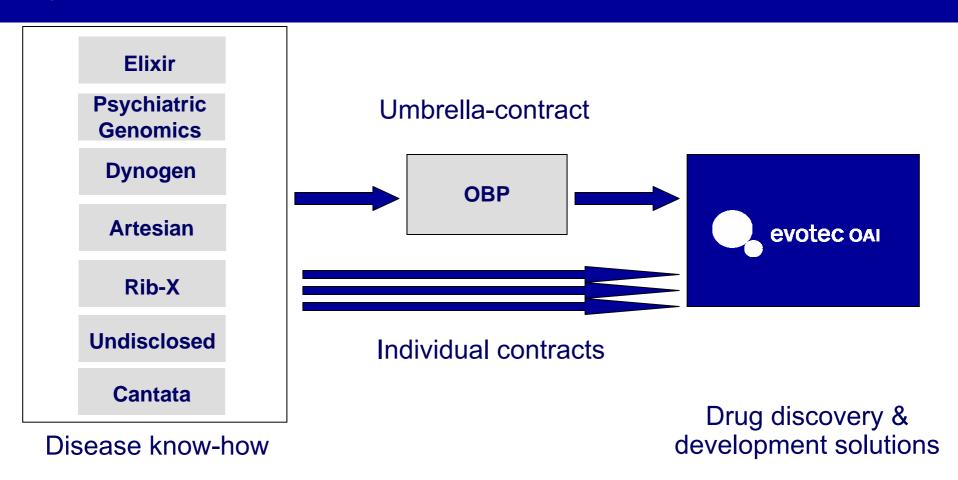
- Partnership started 1997 with joint development of EVOscreen® uHTS technology
- Recent deal extends into assay development and cellular assay screening

Roche

- Partnership initiated in 2001 in high quality chemical library synthesis
- Recent expansion of relationship into medicinal chemistry on a high profile oncology compound



VC firm OBP: 7 deals with portfolio companies > 20 FTEs involved



⇒ We recently closed a comparable deal with MPM Capital



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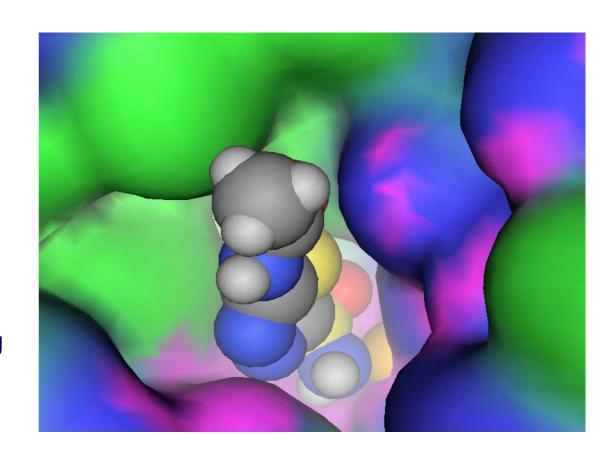
Summary of Discovery and Development Services (DDS)

- Revenues 2003: EUR 61.2m (+4%), Headcount: 440
- Product development
 - Integration of random and rational discovery progressed
 - X-ray crystallography group added
 - New ADMET assays handed over to operations
 - Informatics tools provide quick decisions
- Capacity expansion in Abingdon completed
- Improved integration of formulation capabilities at ProPharma



Increased use of rational design approaches: Virtual screening

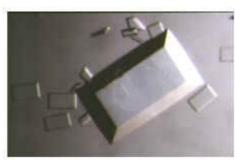
- Docking performed using GOLD
- Grid of up to 250 PCs and a Linux cluster
- Database of 5.5 million compounds
- Established success in finding hits for enzyme targets

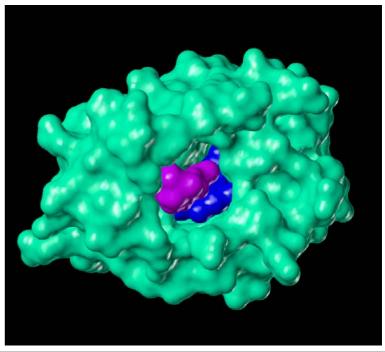




Increased use of rational design approaches: Structural biology

- Established capability in protein engineering and purification for crystallography studies
- Experienced X-ray crystallography team recruited
- Rapid crystallisation screening
- De novo protein structure determination
- Structure determination of ligandprotein complexes







Structural biology facilities



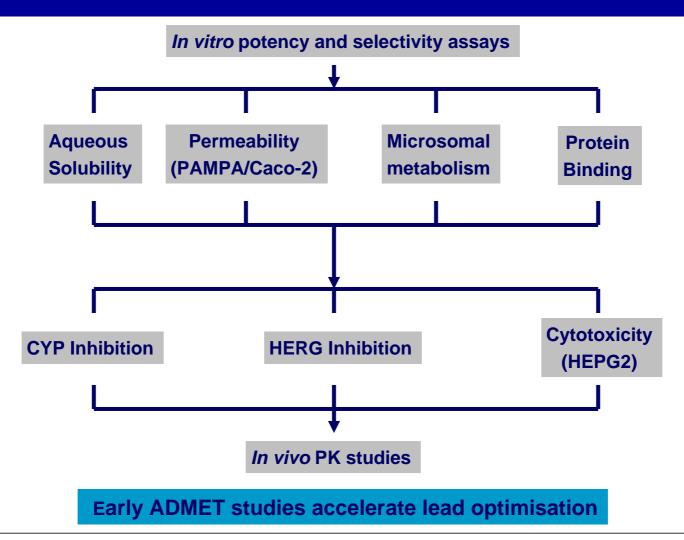
- Automated crystallisation facilities
- Fully equipped X-ray crystallography laboratory
- Access to synchrotron radiation sources for HT data collection
- Ideally placed to exploit Diamond, the third generation Synchrotron site in Oxford

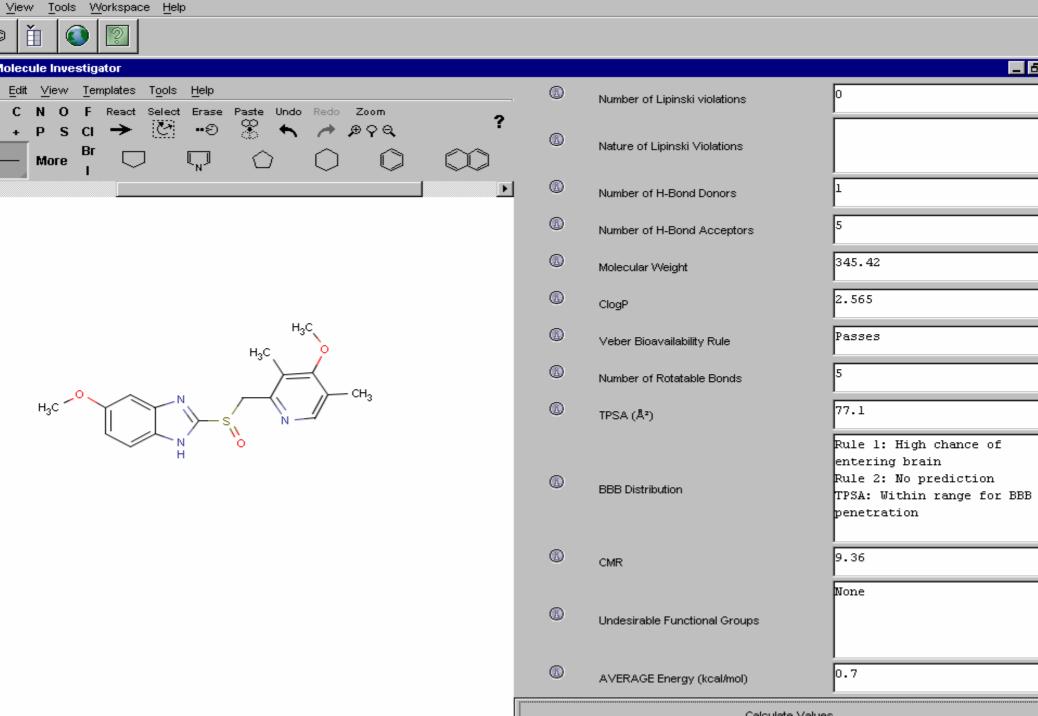






Full in vitro ADMET service now available







Discovery capacity extension in Oxford: Building 114 – space for 200 staff





ProPharma: Sophisticated drug formulation development and manufacture

- Parenteral formulation development
 - Pre-formulation
 - Formulation
 - Freeze-drying
- Sterile / aseptic manufacture (including cytotoxics)
- Labelling for clinical trials
- Regulatory support





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Discovery Programs Division (DPD): Tremendous progress

- Revenues 2003: EUR 1.5m, Headcount: 28
- Business model:
 - Create internal discovery of drug candidates for out-licensing
 - Long-term upside through more significant milestones and royalties
 - Leverage strong DDS discovery and development engine
- Launched research projects in CNS and Metabolic Diseases:
 - Strategic alliance with DeveloGen in Metabolic Diseases

 - ENS licensed 5 CNS compounds from Roche
- ENS prepared for venture capital financing



Evotec OAI / DeveloGen:Discovery programme in obesity/diabetes/metabolic syndrome



Disease know-how: 200 primary, 30 validated targets Pharmacology



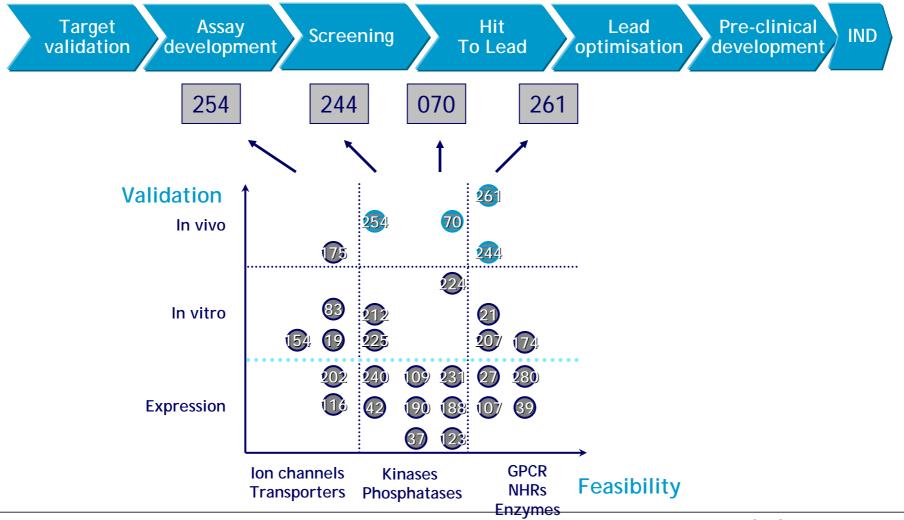
Discovery and development:
Assay development
Screening
Medicinal chemistry
Drug manufacture

Leads, POCDs, INDs, Phase I / IIa candidates

- Top expertise and professional resources over the entire value chain
- Rewards and cost to be shared 50:50
- > 40 scientists
- Early alliance with pharma partner

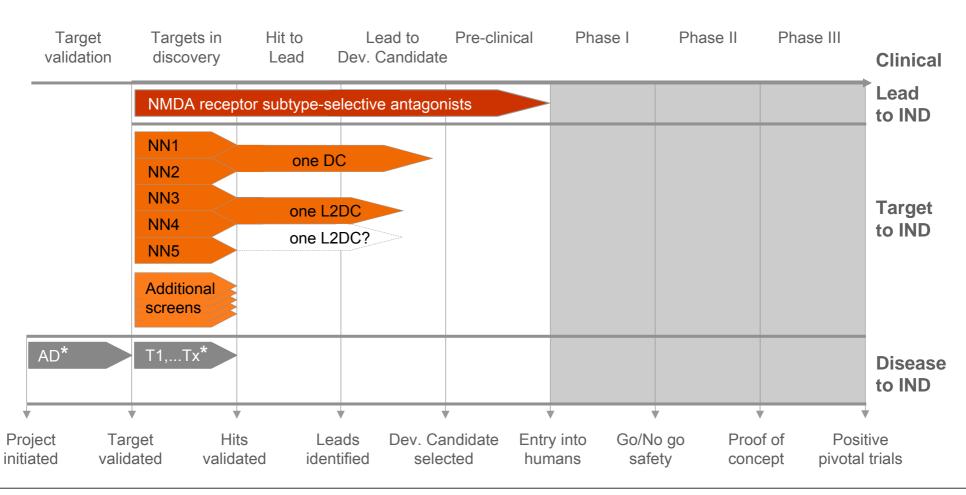


The DeveloGen / Evotec OAI JV's expanding project portfolio





Projected ENS discovery portfolio 2004





NMDA receptor NR2B subtype selective antagonists: Background

- Extensive studies over the last 15 years indicate a role for NMDA receptor antagonists in the treatment of neuropathic pain, Parkinson's disease, Alzheimer's disease, stroke and traumatic brain injury
- However, the clinical development of non-selective antagonists has been hampered by a low therapeutic window due to mechanism related, CNS side-effects
- Memantine, a low affinity, non-selective NMDA receptor antagonist recently approved in Europe and the US for treatment of moderate-to-severe AD
- NMDA receptor subtypes exist which contain different NR2(A-D) subunits
- Compounds highly selective for NR2B subunit-containing receptors have been identified that retain many of the beneficial effects of non-selective compounds but show much improved side effect profiles



NMDA receptor NR2B subtype selective antagonists: Key features

- Portfolio of six patents covering 5 lead compounds plus further compounds protected by these patents
- High affinity and selective compounds with good CNS bioavailability and high potency in vivo, including orally
- Lack HERG channel blocking activities (QT-prolongation) and activities at other known receptors and ion channels
- Active in pre-clinical studies of neuropathic pain and Parkinson's disease and neuroprotection (e.g. stroke) with large therapeutic window with respect to adverse side-effects



Target-to-IND programme: Lead discovery status

	Status
Target 1	uHTS completed, hits
Target 2	uHTS completed,
	Hit2Lead phase
Target 3	uHTS assay
Target 4	Assay development
Target 5	Assay development



Disease-to-IND programme: ENS Alzheimer's Disease programme

 Phase A: Identification of target and marker candidates for AD using differential gene expression analysis on human brain

tissue samples and integration with clinical genetics data

Phase B: Validation of targets and markers in human primary

tissues, cellular and animal disease models

(ongoing)

 Phase C-E: Identification of leads, chemical validation in animal disease models, and development to drug candidates for AD





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Tools and Technologies: Strong year for Evotec Technologies

- Revenues 2003: EUR 18.7m (+64.0%), Headcount: 84
- Quality provider of life science tools and technologies
- First full year as stand alone company
- Products:
 - Two new bench-top devices launched: Elektra and Clarina II
 - Embargo for EVOscreen® is coming to an end
 - Successful collaborations with Pfizer and Olympus continued
- Large number of customer relationships secured
- Pfizer became 10% shareholder in Evotec Technologies
- US base established



Evotec Technologies: A reliable technology partner for critical life science tools and applications





EVOscreen®:An industry standard for uHTS

The embargo with its technology partners is coming to an end



NanoScreening: Savings & Quality

	HTS (50 µl, 384w-plate)	NanoScreen (1.2 µl, 2080w-plate)
# of plates	1421	284
Reaction mix (incl. setup; 'safety' factor)	27.3 Liters (40.9 Liters; f= 1.5)	709 ml (850 ml; f= 1.2)
Compound (MW 500, 10 μM)	250 ng	6 ng
Enzyme (20 nM, 80 kDa)	44 mg	1.14 mg
Substrate amount (100 μM, MW 1000)	2.73 g	71 mg
Substrate cost (1 mg 200 \$)	546'000 \$	14'200 \$

Example: protease assay with fluorescence read-out.

Library size 500'000 compounds, control wells on each plate, 352 compounds on 384w-plate, 1760 compounds on 2080w-plate. In addition, excellent sensitivity for the confocal detection setup with single-molecule detection in NanoScreenian will allow even lower substrate concentrations.

Additional Benefits: - low-affinity protein-peptide interactions possible (K_D>1μM)
- single turnover-enzymology possible (ultimate sensitivity)

Time!

Costs!

Quality!

DDT Boston, Aug. 4-9, 2002, Slide 29

U NOVARTIS -









evotec oai



Clarina™ II workstation: uHTS Evoscreen® technology in bench-top format



- Same cutting-edge technology as in EVOscreen®
 - Data quality
 - Flexibility: 2080 to 96 well plates
 - Easy to use
- Seamless technology solution:
 - Assay development
 - uHTS
 - Secondary screening
- Valuable tool for:
 - Academia
 - Open access labs in pharma
 - Biotech companies
 - Current customer base



Elektra: Addressing a critical bottleneck in cell biology

Generation of positive clones

Duration	Standard	Elektra
Process Step		
Transfection	1 h	1 h
Transient expression	2 d	2 d
Selection	14 d	7 d
Cloning		
Identify&pick clones	1 d	1 d
Amplify clones	14 d	
Test clones	2 d	
Re-cloning		
Identify&pick clones	1 d	
Amplify clones	14 d	
Test clones	2 d	
	> 7 weeks	10 days





Evotec Technologies in the US

Since 01/2003: Evotec Technologies into Technology Center Hialeah



08/2003: Foundation of

Evotec Technologies, Inc.

EIN 20-0078154 (a Delaware Corporation)

- ET's current US presence:
 - Sales managers on East Coast
 - Exclusive agent on West Coast
 - Application lab in Florida for training, demos and spare parts
 - Experienced German field service engineer to build up own service team
- Boston site to be in operations by mid2004

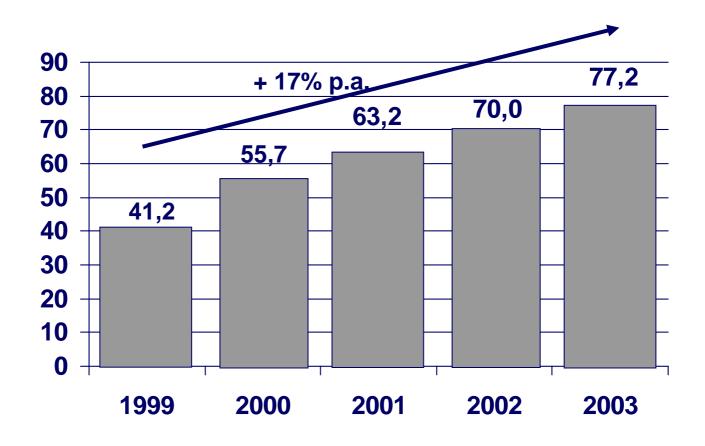


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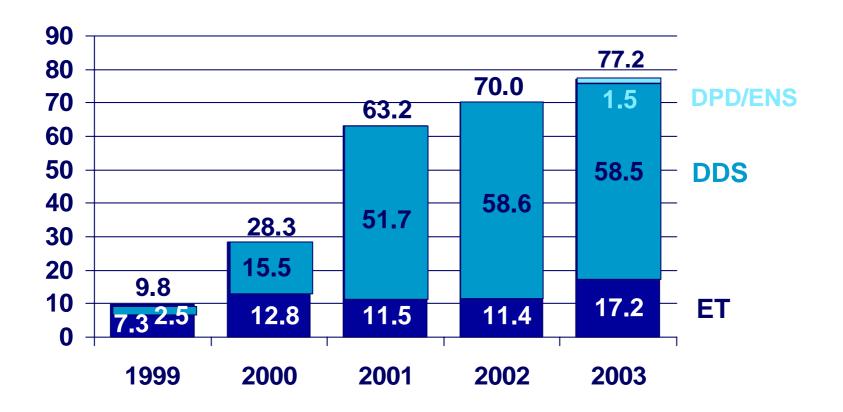


Achieved revenue target, 4 year proforma CAGR of 17%



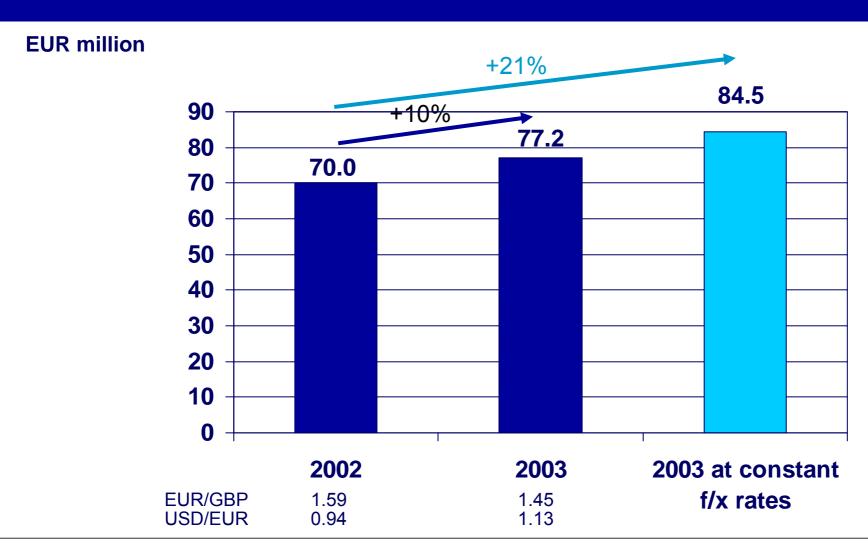


Evotec Technologies contributed strongly to revenue growth





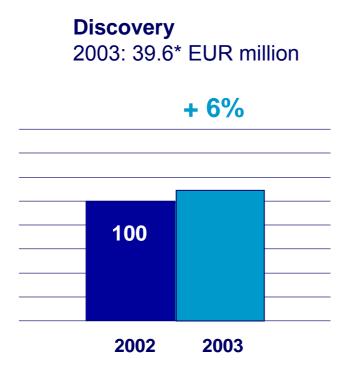
At constant currencies, we would have grown 21%

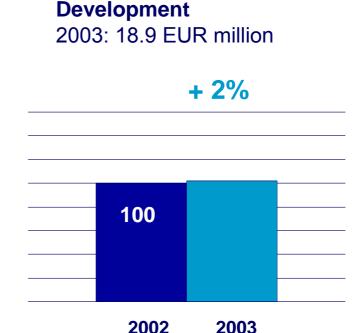




Revenue by product group of DDS: Despite f/x effects, growth in both product lines

(indexed)



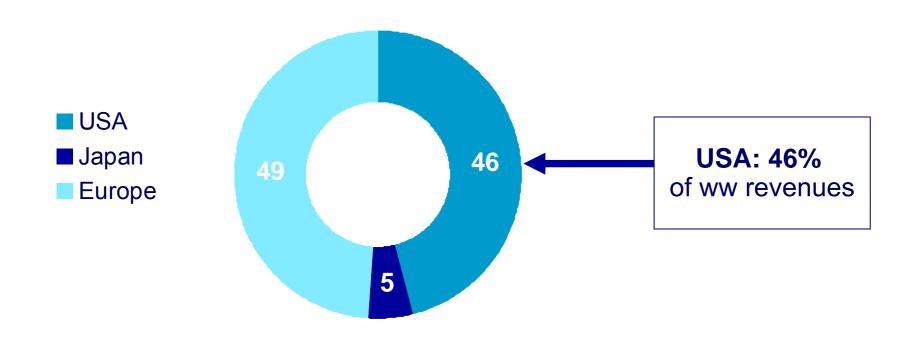


^{*} incl. Intra-group revenues



Revenue by regions: Continued high weight of US business, despite USD decline

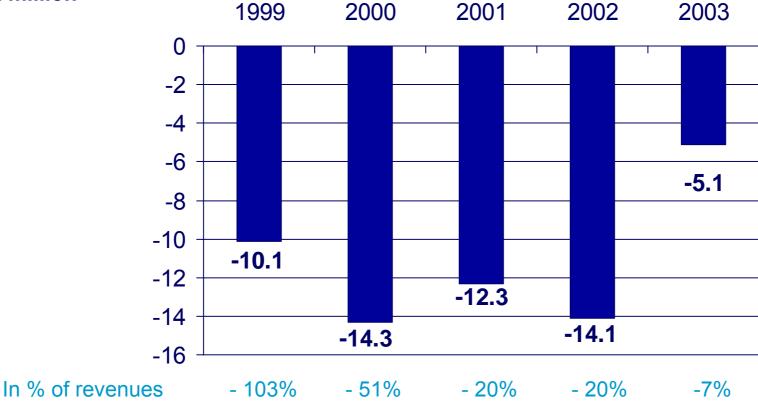
in % of revenues in EUR





Operating result*: Cost reductions resulted in significant improvement

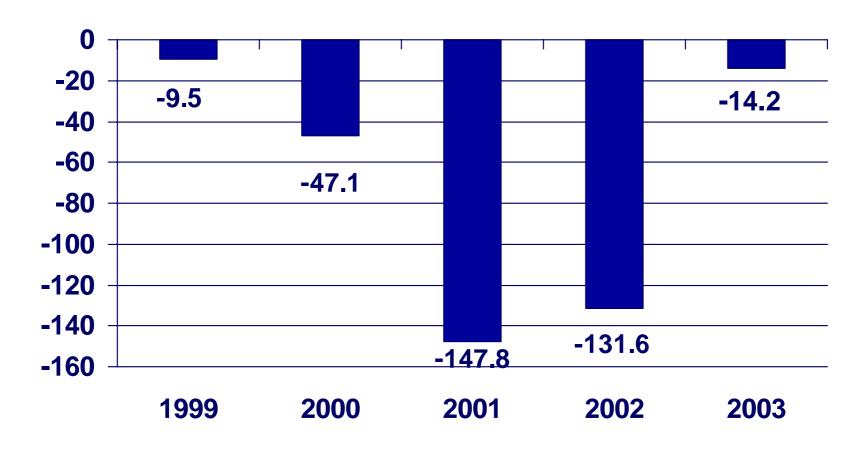




^{*} before amortisation and impairment



Net income: Improvement supported by absence of goodwill impairment



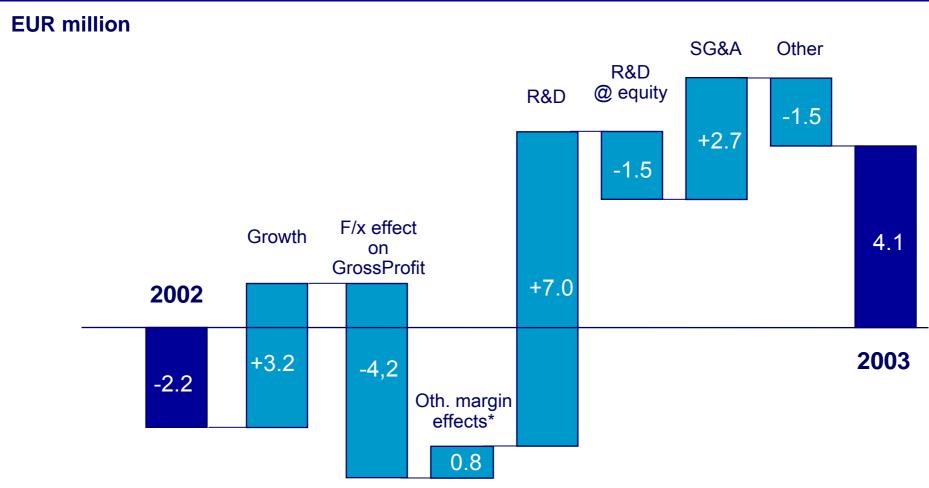


EBITDA: Positive for the first time in company history





EBITDA effects: Improvement primarily from SG&A and R&D reduction



^{*}Sales mix, utilisation/efficiency, pricing

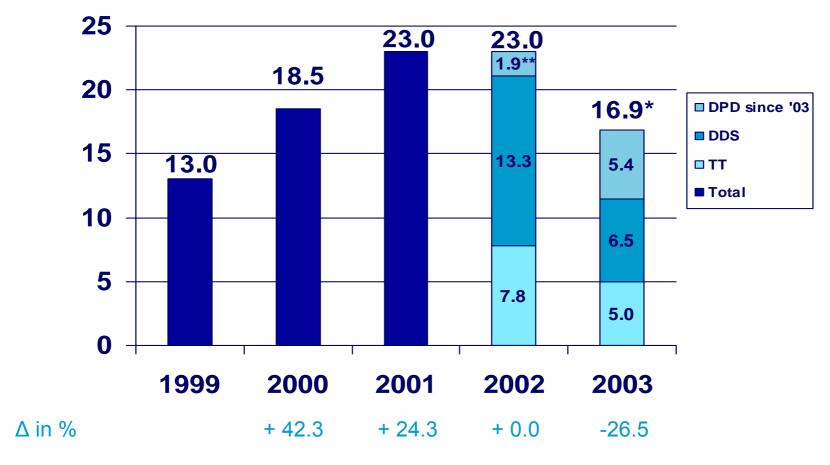


Group P & L

EURO million	2003 Actual	2002 Actual	% vs. Act 02	
Revenues	77,2	70,0	+10%	Delivering on promises
COGS	-46,2	-38,5	-20%	
Gross Profit	31,0	31,5	-2%	
Gross Profit in % of Sales	40,2%	45,0%		US Dollar impact
R&D	-15,5	-23,0	+33%	Reduction of technology R&D
SG+A	-17,9	-20,5	+13%	37
Other Operating Exp.	-2,7	-2,1	-29%	
Amortization Intangibles	-10,7	-121,4	+91%	
Operating Income	-15,8	-135,5	+88%	
Equity in net loss of affiliate	-1,6	-0,1		DeveloGen JV
EBIT	-16,9	-134,7	+87%	
Net Income	-14,2	-131,6	+89%	Absence of goodwill impairment
EBITA	-6,2	-13,3	+53%	+ cost containment
Depreciation	-10,3	-11,1	+7%	
EBITDA	4,1	-2,2	+286%	Positive



Reductions in platform R&D (DDS + TT), growing emphasis on discovery R&D (DPD)

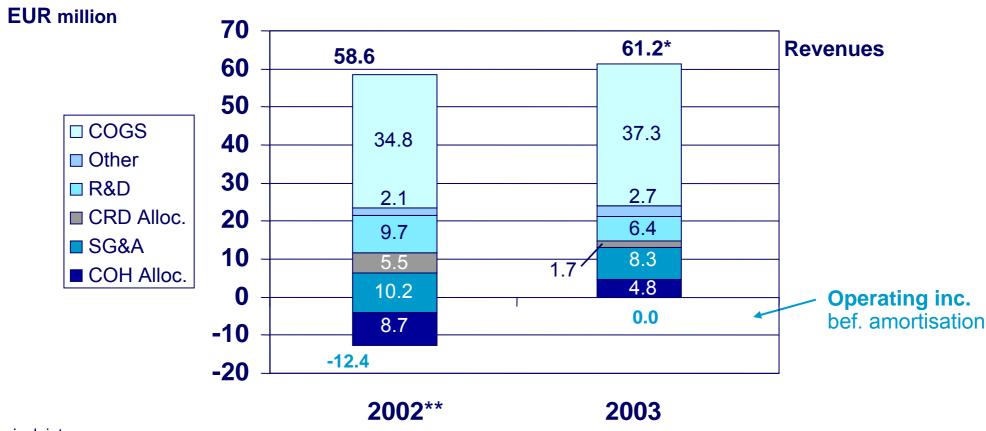


^{*} Incl. €m 1.4 R&D expenses shown at equity, ** ENS GmbH



DDS has achieved positive operating income before amortisation

P&L Discovery and Development Services (Segment)



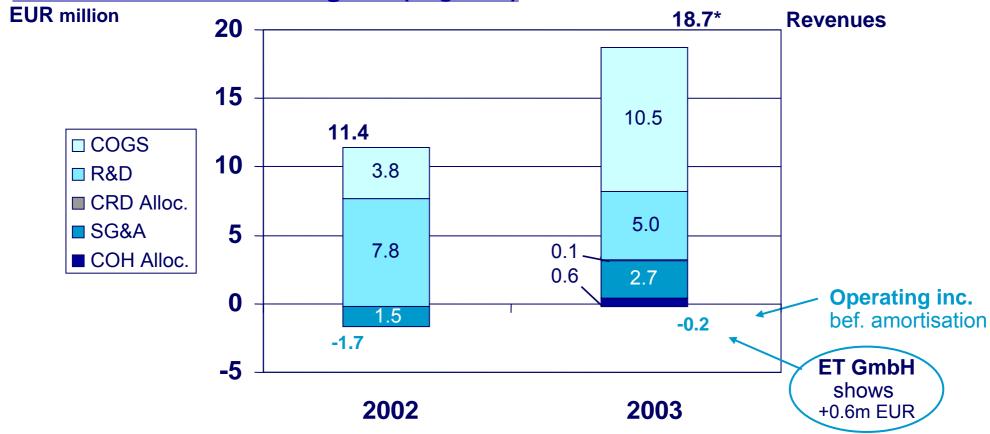
^{*} incl. intragroup revenues Actual '03: 2.6m

^{**} incl. early DPD activities



Evotec Technologies streamlined operations and came close to positive operating result before amortisation





*incl. intragroup revenues Actual '03: 1.5m



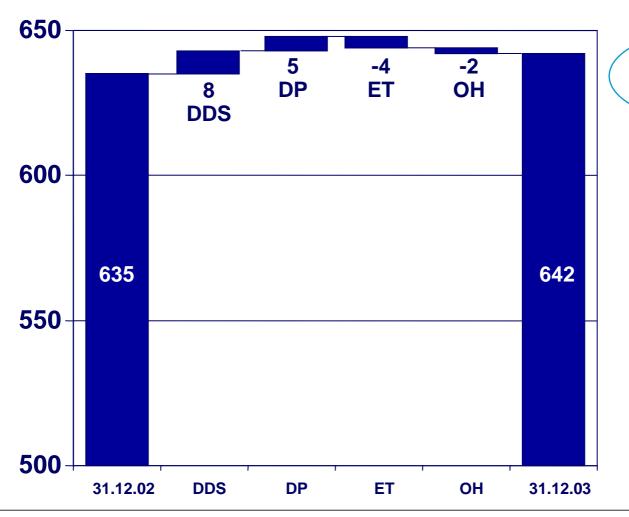
Break even in Operating Result in established businesses, supported creation of Programs Division with > EUR 5 m

EURO million	DDS Actual	Discovery Programs Actual	ET Actual	Consol Actual	Group Actual
Revenues Intra-Group Revenues	58,6 2,6	1,5 0,0	17,2 1,5	0,0 -4,1	77,2 0,0
Total CoGS	-37,3	-0,7	-10,5	2,2	-46,2
Gross Profit G.P. in % of revenues	23,9 39,0%	0,8 55,6%	8,2 43,7%	-1,9	31,0 40,2%
R&D SG+A Other Operating Expenses	-8,1 -13,1 -2,7	-4,3 -1,8 0,0	-5,1 -3,3 0,0	2,0 0,2 0,0	-15,5 -17,9 -2,7
Operating Income before amortisation and impairs	0,0 nent	-5,3	-0,2	0,4	-5,2



Continued hiring in drug discovery

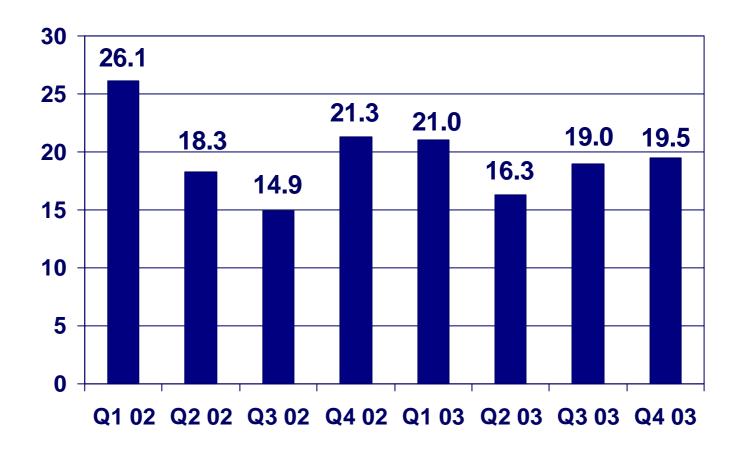
Headcount



Net headcount increase: +7 employees

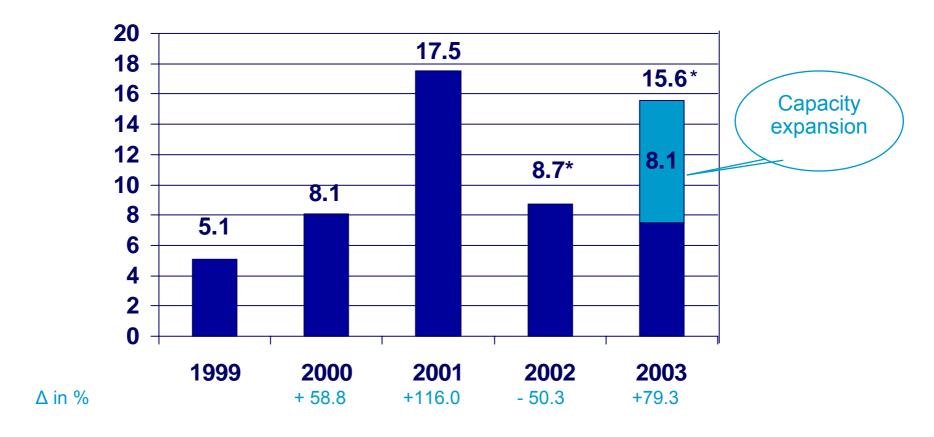


Cash and marketable securities: We have successfully managed our cash position





Capital expenditures: Significant investment in new labs on top of regular capex



^{*}including capital leases



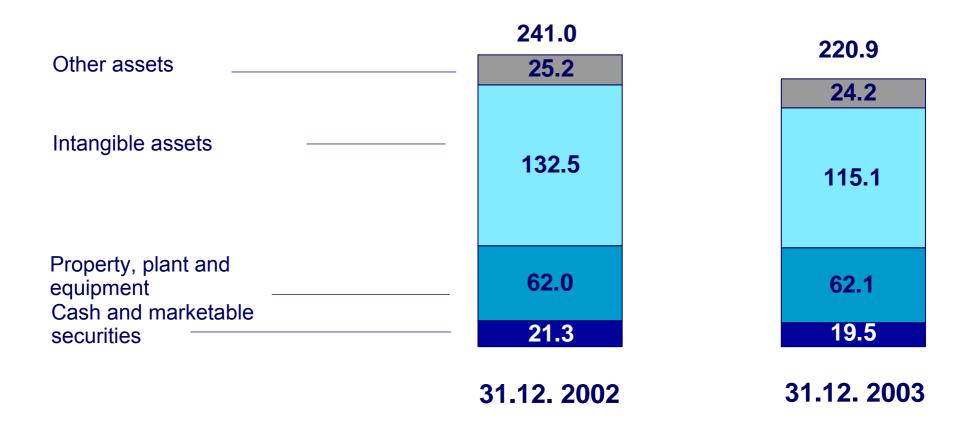
Cash Flow: Strong operating CF helped finance our investments

EURO million	31.12.03 Actual	31.12.02 Actual	% vs. Act. 02
Net cash from operations	+7,8	-1,0	+881%
Net cash from investing** thereof leasehold improvement	-15,6 -8,1	-7,0 -2,1	-123%
Net cash from financing** thereof loan finance thereof capital increases subsidiaries	+7,2 +5,5 +3,0 *	+4,1 +4,1	+75%
Net in-/decrease in cash and securities	-0,6	-3,9	+85%
F/X Differences	-1,2	-2,7	+55%
Cash and securities at beginning of period Cash and securities at end of period	+21,3 +19,5	+27,8 +21,3	-23% -8%
working capital movement	-2,3	0,1	

^{*} Pfizer capital contribution

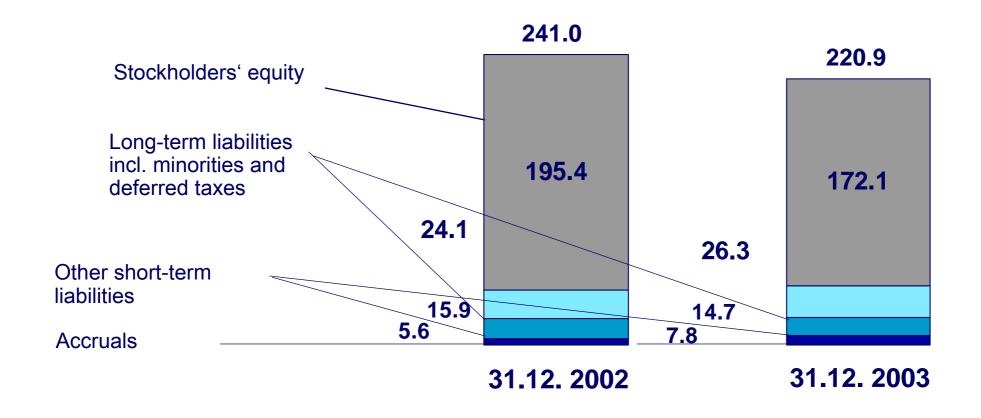


Assets: Regular amortisation continued to reduce our intangibles





Liabilities and stockholders' equity: Solid equity position, stable long-term liabilities





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2004 expectations

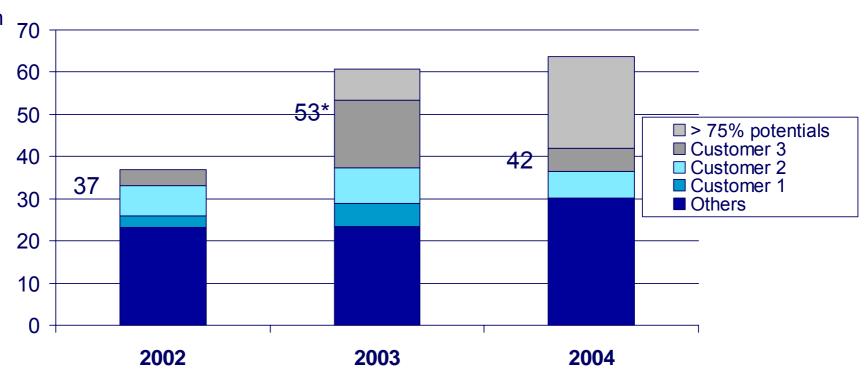
- Market weakness appears to have bottomed out at the end of 2003.
- Signs of recovery in core markets
 - US and Japan
 - Europe lagging behind
- More significant growth only in the second half of 2004
 - Time lag between funding events and orders placed
 - Unfavourable foreign exchange rate Euro / Dollar 1st half 2004 / 2003
- Continued growth and good operational performance for full year 2004
 - Individual quarters will deviate from last year's performance (EVOscreen)
- Ideally positioned to benefit from upswing in the outsourcing market
- Growth in internal programmes



Good order book and prospects

Sales and order book

Status as of February each year



^{*} Adjusted for a contract change with customer 2 in 2003



Synergistic partnerships

