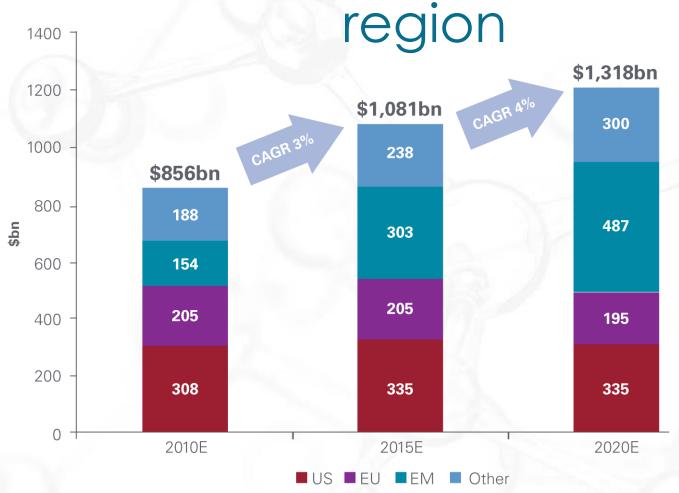
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PHARMACEUTICAL PROSPECTS: A SECTOR & COMPANY OVERVIEW





Pharma sales growth to 2020 by



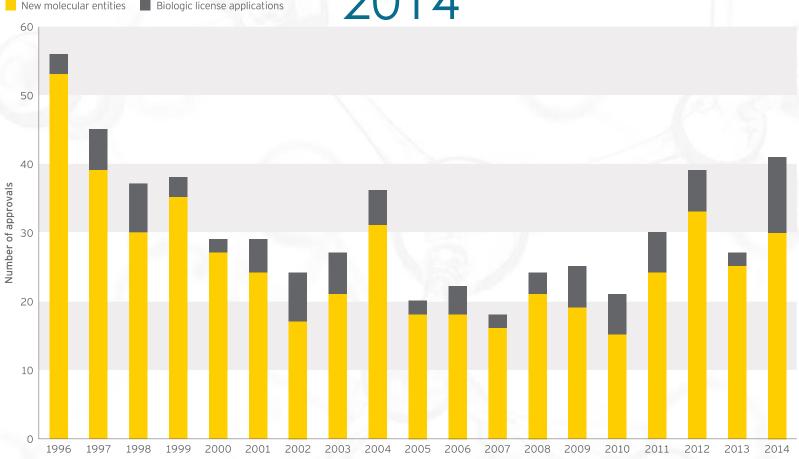
New product Approvals Forecast to Average 35 Per year

>900 products
In the pipeline
2/3 of which are
Small
molecules

Source: IMS, May 2011 & KPMG, Oct 2011



FDA product approvals, 1996— New molecular entities Biologic license applications 2014



Source: EY, Beyond borders report, 2015



Small Molecule API Trends

Small Molecules Continue to Drive Pharmaceutical Growth:

- ► Accounting for 82% of NDA applications in 2014
- ▶ 60% of New Chemical Entities
- Can be engineered to deliver small dose effect
 - Economic advantage
 - More amenable to oral formulation
 - ► Clinical trials can be simpler
 - ▶ PR&D can be cheaper
 - ► Shipment and storage conditions can be easier



Small Molecule API Trends

Small Molecules Continue to Drive Pharmaceutical Growth:

- ▶ Targeted therapeutic trends
- Higher molecular complexity / higher potency
- ➤ Smaller batch size
- ▶ Projection that in 15 years the ratio of NCE / BIO could still be 60/40



Small Molecule API Trends

Number of Small-Molecule Drugs, 2013 and Projected for 2010

Revenues	2013	2020
Over \$1 billion	75	89
\$500 million to \$1 billion	111	140
\$250 million to \$500 million	160	227
\$100 million to \$250 million	345	407
\$50 million to \$100 million	328	409
Less than \$50 million	1991	2180
TOTALS	3005	3552



North East Pharma Sector

Strong and Supportive Capability:

- >200 Life Science & Healthcare Companies
- ► Turnover > £10billion
- ➤ 33% Of UK Pharma GDP
- ► Full Range of Capabilities
 - Preclinical and clinical development
 - Clinical trial management
 - Drug Substance and Drug Product manufacturing



North East Pharma Sector

Strong and Supportive Capability:

- ► Strong CMO capacity
- ► Strong University Presence
- ► CPI / National Biologics Centre / National Formulations Centre



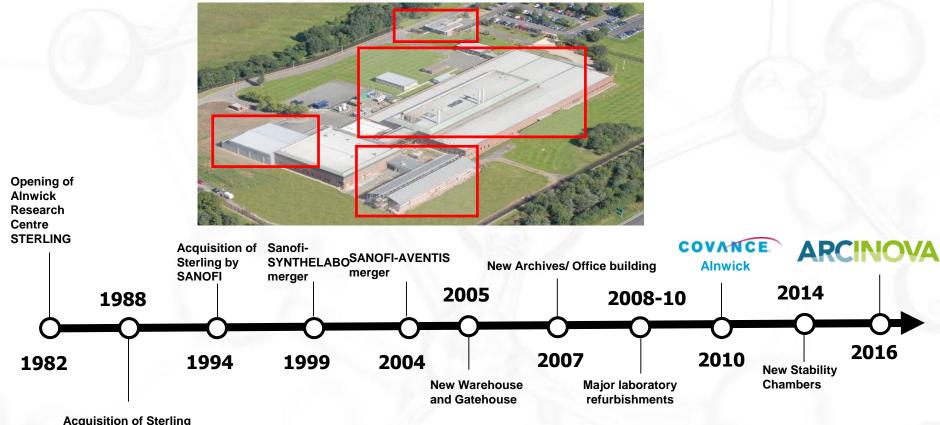
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- ► Site area : ~12.8ha (1 ha = 10,000 m²)
- ► Total foot print of site buildings ~10,000 m²
 - Of which 2/3 is laboratory space





Well-equipped site & rich history

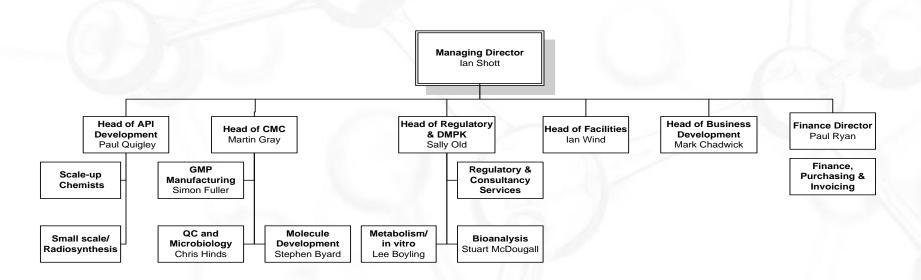


Acquisition of Sterling by EASTMAN-KODAK

- ▶ 60 current customers worldwide
- ► Biotechs, speciality pharmas, large pharmas



Experienced Team



- Over 50 project staff (chemists, analysts, engineers)
- ► Average 20 years industry experience



Small Molecule Capacity

Key Differentiators:

- Strong and established route scouting capabilities
- Established chemistry base
- Extensive Drug Substance / Drug Product development expertise
- Established API analytical capabilities (chemical and physical)
- ▶ GMP compliant site
- Well established infrastructure



Investment - Capability

Building:

- ► Strong Process Development Expertise
- ► Range of API development and manufacturing scales
- ► Alliance partners for further scale up
- ► Ability to handle multiple API classes
 - Standard
 - Radiolabelled
 - High toxicity / Potency
- ► Flexibility to introduce new technologies:
 - Synthetic Biology
 - Continuous processing
- ▶ Nimbleness, agility and innovative service capabilities



Investment - Infrastucture

- ▶ Driving investment from 5lt to 80lt scale
- ► Integrating with preferred scale up partners
- ► Tying into design capabilities of established partners (BPE)





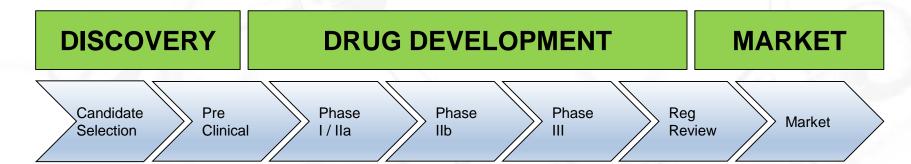








Offerings



Contract Research Services:

Capability for the development of small molecule drug candidates from discovery through pre-clinical and clinical phase I, II and III to launch. These services can also be applied to a range of life sciences applications.

Contract Development Services:

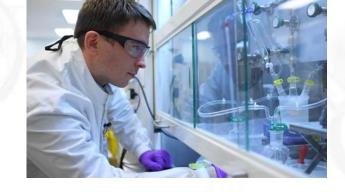
Capability to include Process Research, Development and Scale-Up, drug candidates in both Development and launched to the Market, at relatively small scale using synthetic biology and chemistry.



Offerings

Efficient, end to end solutions, including:

- ▶ Drug substance synthesis & support intermediates & APIs
- ▶ Isotope synthesis/radiochemistry
- Drug product formulation
- ► High potency/high hazard materials



► Full bioanalytical support from pre-clinical through clinical trials



Compliance

- ▶ 10 successful MHRA GCP/GLP inspections since 2000
- ▶ 10 successful MHRA/FDA GMP inspections since 2005





Summary - What makes us different?

- ► Fully integrated provider
 - ► Pre-clinical through to small scale commercial
 - ► Cohesive API development, CMC, BioA & metabolism studies
- ▶ Big company track record/compliance
 - ➤ 34 years successful delivery of projects
- Speed and agility of a small company
 - ► Responsive team, track record of delivery
- ▶ World class facilities and capabilities
 - Modern state of the art research centre



Strategy

Market segmentation to achieve four key target **customer groups**:

- ► Emerging Pharma / Biotechs seeking a full service partner
- Midsized pharmaceutical companies seeking strategic suppliers
- ► Global Pharma seeking trusted low risk and cost effective preferred suppliers of carefully targeted services
- ► Innovative Life Science companies and organisations (e.g. CPI) requiring professional consultancy and critical service support



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Thank you

