

<u>Connection details</u> https://astrazeneca.zoom.us/s/99546038702 +44 203 481 5237 +46 8 5052 0017 +1 301 715 8592 ID: 995 4603 8702 | PW: 12121220

Accelerating AstraZeneca's strategic and financial development

AstraZeneca to acquire Alexion

12 December 2020

This webinar and conference call for investors and analysts is being recorded



Forward-looking statements

This document may include statements that are or may be deemed to be forward-looking statements. These forward-looking statements may be identified by the use of forward-looking terminology, including the terms "believes", "estimates", "envisages", "plans", "projects", "anticipates", "targets", "aims", "expects", "intends", "may", "will" or "should" or, in each case, their negative or other variations or comparable terminology, or by discussions of strategy, plans, objectives, goals, future events or intentions and include, but are not limited to the ability of the parties to consummate the proposed acquisition on a timely basis or at all, the ability of the parties to satisfy the conditions precedent to consummation of the proposed acquisition, including the ability to secure the required regulatory approvals on the terms expected, at all or in a timely manner, the ability of AstraZeneca to successfully integrate Alexion's operations, and the ability of AstraZeneca to implement its plans, forecasts and other expectations with respect to Alexion's business after the completion of the proposed acquisition and realise expected synergies. Economic, competitive, governmental, technological and other factors that may affect AstraZeneca's and Alexion's operations are discussed in the section entitled "Risk Factors," in each of AstraZeneca's Annual Report on Form 20-F for the year ended 31 December 2019, and Alexion's Annual Report on Form 10-K for the year ended 31 December 2019, in each case as amended by any subsequent filings made with the SEC. These forward-looking statements include all matters that are not historical facts and involve predictions. Forward-looking statements may and often do differ materially from actual results. Any forward-looking statements reflect AstraZeneca's and Alexion's current views with respect to future events and are subject to risks relating to future events and other risks, uncertainties and assumptions relating to AstraZeneca's or Alexion's results of operations, financial position, liquidity, prospects, growth or strategies and the industries in which they operate. Forward-looking statements speak only as of the date they are made and cannot be relied upon as a guide to future performance. Save as required by law or regulation, AstraZeneca and Alexion disclaim any obligation or undertaking to release publicly any updates or revisions to any forward-looking statements in this document that may occur due to any change in their expectations or to reflect events or circumstances after the date of this document.

Nothing in this document should be construed as a profit estimate or profit forecast. No statement in this document, including statements that the acquisition is accretive to core EPS or enhancing to core operating margins, should be interpreted to mean that earnings per share of AstraZeneca or Alexion for the current or future financial years would necessarily match or exceed the historical published earnings per share or core operating margins of AstraZeneca or Alexion.

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In connection with the proposed acquisition, AstraZeneca intends to file a registration statement on Form F-4 with the SEC, which will include a document that serves as a prospectus of AstraZeneca and a proxy statement of Alexion (the "proxy statement/prospectus"), Alexion intends to file a proxy statement with the SEC (the "proxy statement") and each party will file other documents regarding the proposed acquisition with the SEC. Investors and security holders of Alexion are urged to carefully read the entire registration statement and proxy statement/prospectus or proxy statement and other relevant documents filed with the SEC when they become available because they will contain important information. A proxy statement/prospectus or a proxy statement will be sent to Alexion's shareholders. Investors and security holders will be able to obtain the registration statement and the proxy statement/prospectus or the proxy statement free of charge from the SEC's website or from AstraZeneca or Alexion as described in the paragraphs below.

Participants in the solicitation

Alexion and certain of its directors, executive officers and employees may be deemed participants in the solicitation of proxies from Alexion shareholders in connection with the proposed acquisition. Information regarding the persons who may, under the rules of the SEC, be deemed participants in the solicitation of the shareholders of Alexion in connection with the proposed acquisition, including a description of their direct or indirect interests, by security holdings or otherwise, will be set forth in the proxy statement/prospectus or proxy statement when it is filed with the SEC. Information about the directors and executive officers of Alexion and their ownership of Alexion shares is set forth in the definitive proxy statement for Alexion's 2020 special meeting of shareholders, as previously filed with the SEC on 26 March 2020. Free copies of these documents may be obtained as described in the paragraphs above.



AstraZeneca's transformation has been pipeline-led



Broad presence

Balanced specialty and primary-care franchises¹

Leading emerging markets presence

Strong pipeline

17 Phase III medicines and significant lifecycle projects

Advancing early and mid-stage pipeline

Improving financials

Nine blockbuster medicines

Returned to sustainable revenue and earnings growth

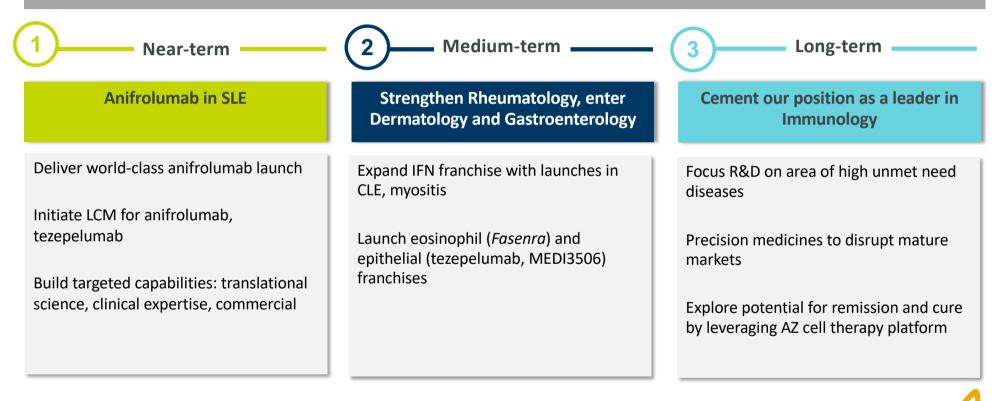
Focus on operating leverage and cash flow

Focused on science and patients Innovative medicines in Oncology and BioPharmaceuticals²

1. In YTD 2020, speciality-care medicines (Oncology, Brilinta, Lokelma, Evrenzo and Fasenra) comprised 53% of total revenue 2. Cardiovascular, Renal & Metabolism and Respiratory & Immunology

AstraZeneca has been expanding its Immunology effort

Focus on achieving leadership in diseases driven by IFN, eosinophilia and epithelial injury across Rheumatology, Dermatology and Gastroenterology



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Alexion is aligned with our strategy Our external growth strategy is anchored in clear principles



Aligns with AstraZeneca's strategy

- Deliver therapy area leadership
- Accelerate innovative science



AstraZeneca adding value through *therapy-area leadership*, *scientific complementarity*, and/or *geographical reach*



Supports top-line growth

Right price, supporting attractive investment return

Immediately accretive to core earnings

Integration manageable; organisation, footprint and culture



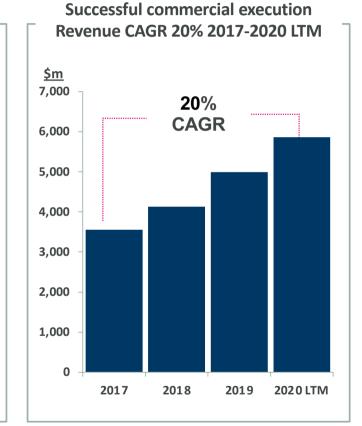
Shared purpose and values

Strong financials

Alexion is a global leader in immune-mediated rare diseases¹

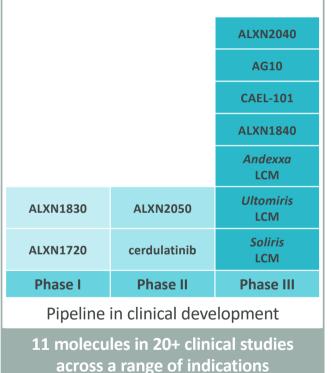
Alexion: market-leading complement franchise

- Track record of success with significant progress made over the last several years (e.g. growth, LCM, pipeline expansion)
- Global leader in the complement system, a critical part of the immune system
- Soliris, the first blockbuster medicine from 2007 for a rare, immune-mediated blood disorder, PNH², followed by aHUS³, gMG⁴ and NMOSD⁵, expanding its use
- Ultomiris, 2nd-generation medicine launched in 2018 (LoE⁶ 2035) with impressive ~70% switch of PNH patients from Soliris⁷ to Ultomiris within 18 months of launch



Source: Alexion financial statements, Bloomberg, FA, LTM as of September 2020

Robust pipeline across immunemediated and other rare diseases



Source: Alexion Capital Markets Day, October 2020; Company information December 2020.



^{1.} A rare disease is a disease impacting less than 200,000 patients in the US $% \left({{{\rm{US}}} \right)$

^{2.} Paroxysmal nocturnal haemoglobinuria 3. Atypical haemolytic uremic syndrome

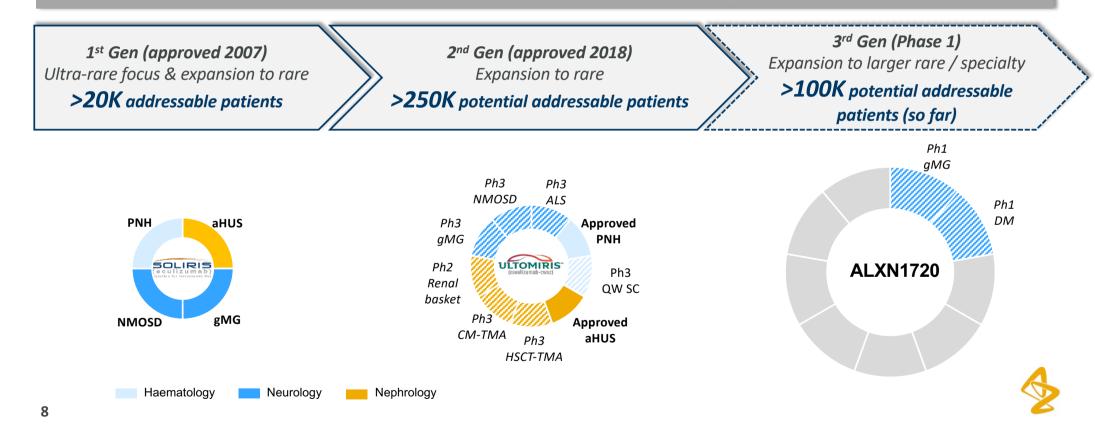
^{4.} Generalised myasthenia gravis 5. Neuromyelitis optica syndrome disorder

⁶. Loss of exclusivity (US) **7**. US, Japan and Germany.

Strong financials

Alexion: a pioneer innovator in C5 complement inhibition Bringing life-saving medicines to more patients

Ultomiris has achieved impressive ~70% conversion in 18 months in PNH across key geographies (US, JP, DE); Ultomiris aHUS conversion underway and tracking PNH



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Strong financials

Alexion: accelerating AZ's strategic and financial development

Compelling scientific complementarity and synergy

- > Increased immunology presence: complement system research platform, currently applied in rare diseases
- > Pipeline further strengthened with 11 molecules¹ across 20+ clinical development programmes
- > Leveraging AstraZeneca's precision medicine capabilities

Combination of two science and patient-centric organisations

- > Focus on science and innovation
- > Patient-centric organisations with high-touch patient support services

Further-sustained, industry-leading revenue growth

- > Attractive growth in specialty and highly specialised / rare disease care
- > Leverage AstraZeneca's global geographical reach to accelerate Alexion's portfolio and pipeline
- > Double-digit revenue growth through 2025

Improved profitability and strengthened cash flow

- > Core operating margin significantly enhanced in the short-term; continued margin expansion. Synergies c.\$500m p.a.²
- > Double-digit Core EPS accretion for the first three years
- > Strong cash flow, rapid debt deleveraging with an ambition to increase the dividend
- > Strong, investment-grade credit rating to provide strategic and financial flexibility

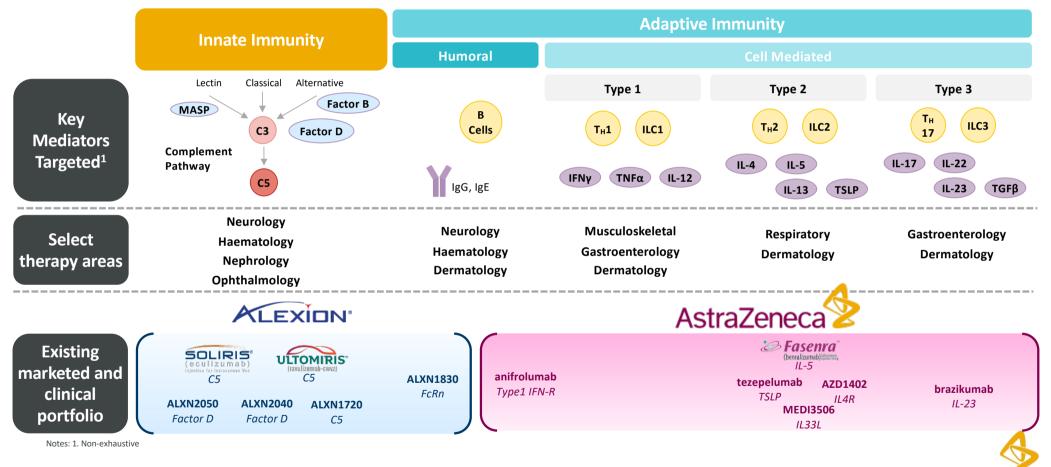


1. See page 7 for details; 2. Synergies are expected from the combined company and are stated on a pre-tax basis with full run-rate expected to be achieved by the end of the third year following completion of the acquisition

Shared purpose and values

Strong financials

Combined R&D capabilities cover both adaptive & innate immunity Improved ability to identify attractive new targets



Shared purpose and values

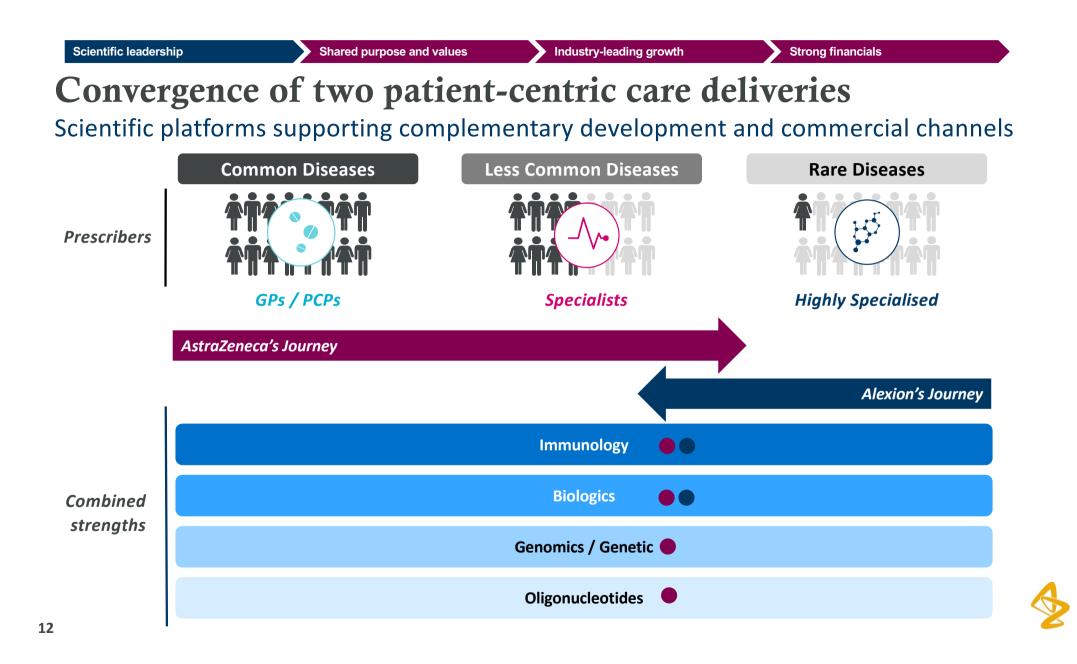
Strong financials

Complement system research platform applicable in many diseases

	Complement tec	hnology platform
Therapeutic areas	Rare diseases	Common diseases
Oncology*	Glioblastoma	Lung Renal Breast Gastric NHL Prostate
Haematology	Cold agglutinin disease PNH TTP <i>wAIHA</i>	
Nephrology	aHUS C3G IgA nephropathy	Haemodialysis
Neurology	GBS gMG NMOSD Dermatomyositis	Multiple sclerosis Mood disorders Dementia Neurotrauma
Respiratory		Asthma Acute respiratory distress syndrome COPD
Cardiovascular	Antiphospholipid syndrome	CAD CHF Myocardial Stroke Diabetic angiopath infarction
Rheumatology	ANCA vasculitis	Rheumatoid arthritis Psoriatic arthritis Osteoarthriti
Dermatology	Pemphigoid Pemphigus Hereditary angioedema	
Ophthalmology	Uveitis	Macular degeneration Glaucoma
Gastroenterology		Crohn's disease Ulcerative colitis
Other	Snake bite Post-infection HUS	Anaphylaxis Transplant Sepsis NAFLD COVID-1 rejection

Abbreviations: aHUS, atypical haemolytic uremic syndrome; ANCA, Anti-neutrophil cytoplasmic antibody-associated vasculitis; C3G, C3 glomerulopathy; CAD, coronary artery disease; CHF, chronic heart failure; COPD, chronic obstructive pulmonary disease; DM, dermatomyositis; GBS, Guillain-Barré syndrome; gMG, generalised myasthenia gravis; HUS, haemolytic uremic syndrome; NAFLD, non-alcoholic fatty liver disease, NHL, non-Hodgkin lymphoma; NMOSD, neuromyelitis optica syndrome disorder; PNH, paroxysmal haemoglobinuria; TTP, thrombotic thrombocytopenia purpura; wAIHA, warm autoimmune haemolytic anaemia. **Bold**: Alexion marketed medicine; *italic*: Alexion pipeline indication. *Chart shows select cancer types.

11 naemogrobinuna; TP, thrombotic thrombotic



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Strong financials

Strong C5 franchise, and a number of innovative assets

Market-leading and highly durable C5 franchise (Soliris / Ultomiris)

- > Proven impressive product conversion expertise; next generation innovative C5 formulations
- > Multi-pronged expansion in neurology driven by Ultomiris (late stage) and ALXN1720 (early stage)
- > Indication expansion into other disease areas with high unmet needs (ALS, HSCT-TMA, CM-TMA, renal basket)

Factor D platform: potential for new Complement franchise

- > Potential for transformational new oral therapy in PNH
- > Significant potential in nephrology, including in areas of high unmet needs (e.g. IgAN, C3G)

Andexxa: strong synergy with AstraZeneca's CVRM presence

- > Compelling clinical profile in an area of high unmet need
- > Large commercial opportunity driven by growing Factor Xa patients

ALXN1840 (Wilson Disease): near-term launch opportunity in an area with significant unmet need

- > Differentiated, competitive clinical profile
- > Potential to be the first new Wilson Disease medicine in decades

CAEL-101: medium-term opportunity in a severe, multi-organ haematological disorder

- > Differentiated, competitive clinical profile with potential to change standard of care
- > Substantial commercial opportunity



Strong financials

Complementary cultures with shared focus on patients

- Two companies with shared mission and purpose to create value for patients through delivery of innovative medicines
 - > Opportunity for shared learnings and innovation across spectrum of primary care, specialty care and rare diseases
 - AstraZeneca: Strong history in primary care with increasing focus on specialty care and precision medicine
 - Alexion: Specialised rare disease field force, high-touch patient services organisation, end-to-end supply chain, innovative technologies for patient-finding and diagnoses critical to success in rare disease
 - > Talented global workforce dedicated to mission-driven performance
 - > Emphasis on building a culture that empowers colleagues and supports long-term success
- Track record for innovation and leadership in respective therapy areas of focus
 - AstraZeneca has a heritage of scientific innovation with first PPI, first EGFRi, first EGFR-790Mi, first PARPi, first immunotherapy for Stage III NSCLC, best-in-class HER2 ADC
 - Alexion is a leader in complement biology and other rare diseases with first C5i, first long-acting C5i, first therapies for HPP and LAL-D, first new therapy for gMG in decades and first NMOSD therapy, first specific Xa inhibitor reversal agents

• Combination creates biopharmaceutical innovator with industry-leading, diverse portfolio

- > Joint history of portfolio and pipeline diversification benefits combined company
- > Opportunity for greater investment in science to develop transformative medicines



¹⁴ Abbreviations: PPI = proton pump inhibitor, EGFR = epidermal growth factor receptor, PARPi = poly-ADP ribose polymerase inhibitors, NSCLC = non-small cell lung cancer, C5i = complement 5 inhibitor, HPP = Hypophosphatasia, LAL-D = Lysosomal acid lipase deficiency, gMG = generalised myasthenia gravis, NMOSD = neuromyelitis optica spectrum disorder, HER2 = epidermal growth factor receptor 2, ADC = antibody drug conjugate

Shared purpose and values

Strong financials

A new pure-play biopharma innovator with strong profitability, and industry-leading science and growth

Broad Presence

Leading commercial infrastructure in NA, Europe, APAC and EMs

Balanced primary care, specialty care and rare diseases franchises

Global R&D platform across multiple centres of excellence

Strong Pipeline

Scientific leadership in Oncology, CVRM and Respiratory & Immunology, with strong expertise in rare diseases

28 Phase III medicines and significant lifecycle projects

Advancing pipeline with breadth and depth of mechanisms

Superior Financials

9 blockbuster medicines¹, growing to 12 by 2023²

Fastest growing global biopharmaceutical company

Sustainable revenue growth and robust earnings

Strong operating leverage and cash flow generation

Combination of two science and patient-centric organisations

Increased operating leverage underpinning strong financial flexibility



1. Defined as product with end-sales in excess of \$1bn globally in 2019. Products include: *Tagrisso, Lynparza, Imfinzi, Farxiga, Calquence, Fasenra, Symbicort, Brilinta, Nexium, Crestor* and *Pulmicort* 2. Defined as product with end-sales in excess of \$1bn globally in 2023. Products include: *Tagrisso, Lynparza, Imfinzi, Farxiga, Calquence, Fasenra, Symbicort, Brilinta, Enhertu, Ultomiris, Soliris* and *Strensiq.* Excludes *Nexium*

Strensia, Kanuma Tagrisso, Imfinzi, Lynparza, Calquence, Kidnev • Soliris LCM⁴, Ultomiris LCM, Andexxa Fnhertu LCM; four other late-stage projects Nine late-stage projects, incl. NMEs² (ALXN1840, CAEL-101, AG10, Liver • • capivasertib. tremelimumab. ALXN2040); four early to mid-stage monalizumab and savolitinib pipeline projects (ALXN1720, Lung • ALXN1830, ALXN2050, cerdulatinib) Mid-stage pipeline, incl. oral SERD, datopotamab deruxtecan, etc. Other immune-mediated diseases

> 3. Revenue last four guarters as of Q3 2020 4. Lifecycle management (new approvals/indications).

Global presence across primary to specialised and highly specialised care

5. Cardiovascular, Renal & Metabolism 6. Total revenue last four quarters as of Q3 2020, includes New CVRM and Respiratory & Immunology, Rest of CVRM and Other medicines excluded 7. Non-alcoholic steatohepatitis.

CVRM⁵, R&I - Organ protection and immune-mediated diseases \$10.0bn | +5%⁶

- Heart Farxiga, Brilinta, Andexxa, pipeline
- Farxiga, Lokelma, Evrenzo, pipeline
- Pipeline in NASH⁷

Symbicort, Fasenra, Breztri/Trixeo, tezepelumab, PT027, early pipeline

Fasenra, tezepelumab, brazikumab anifrolumab

1. Total revenue last four guarters as of Q3 2020 2. New molecular entity.

Oncology

\$10.8bn | +17%1

Lung, ovarian, breast, prostate and

Scientific leadership

blood cancers

Stronger innovative portfolio to drive sustainable growth

Rare diseases

\$5.9bn | +24%³

ALEXION'





Soliris. Ultomiris

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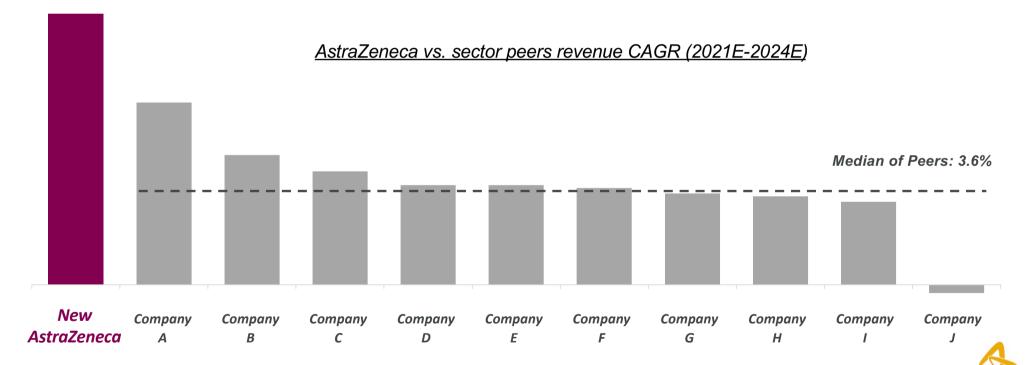
Strong financials

Shared purpose and values

Strong financials

The combination cements AstraZeneca's growth leadership

AZ double-digit revenue growth



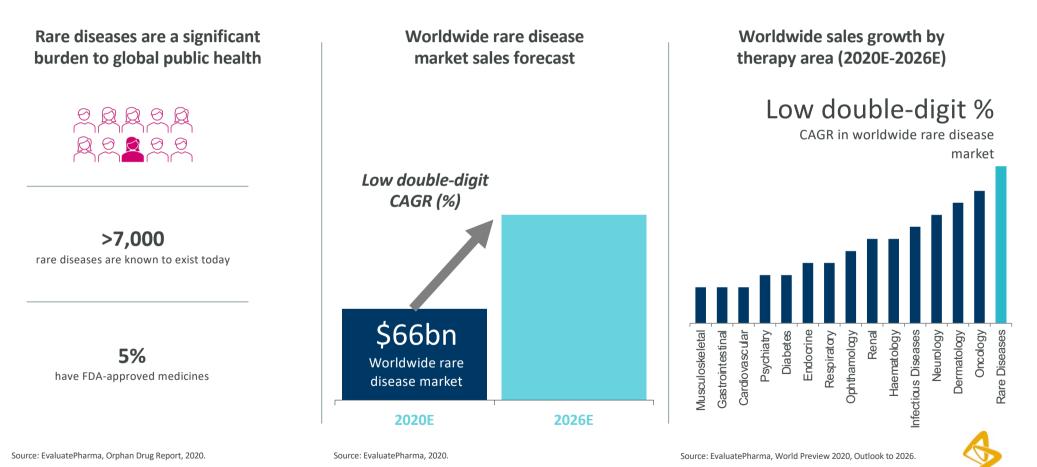
Source: '21-24E CAGR based on FactSet median consensus as of 10 December 2020.

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Shared purpose and values

Strong financials

Rare diseases is a strong growth driver Accelerating growth and innovation in an area of high unmet needs



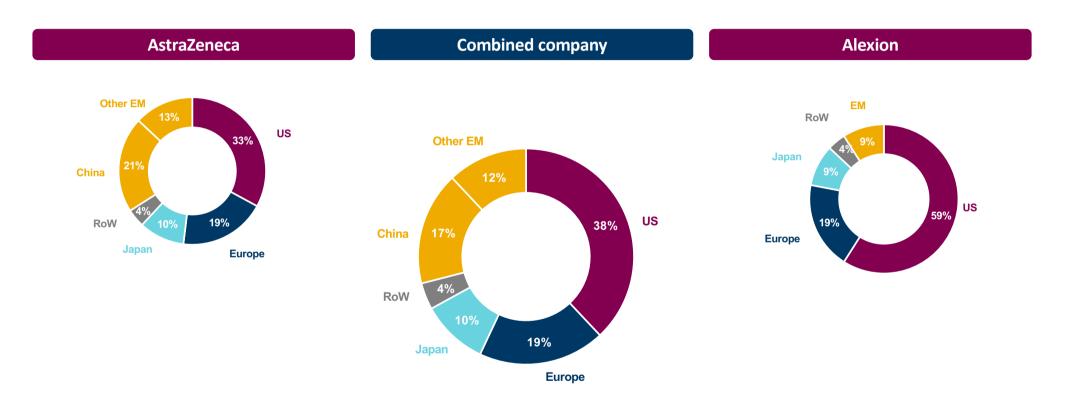
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Strengthening US presence and leveraging global footprint

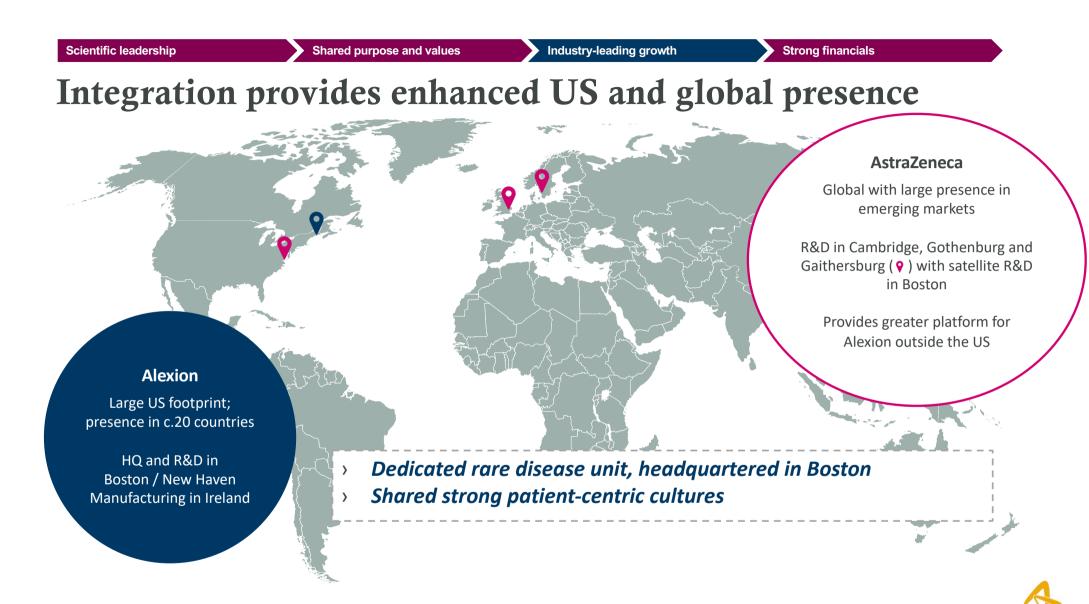
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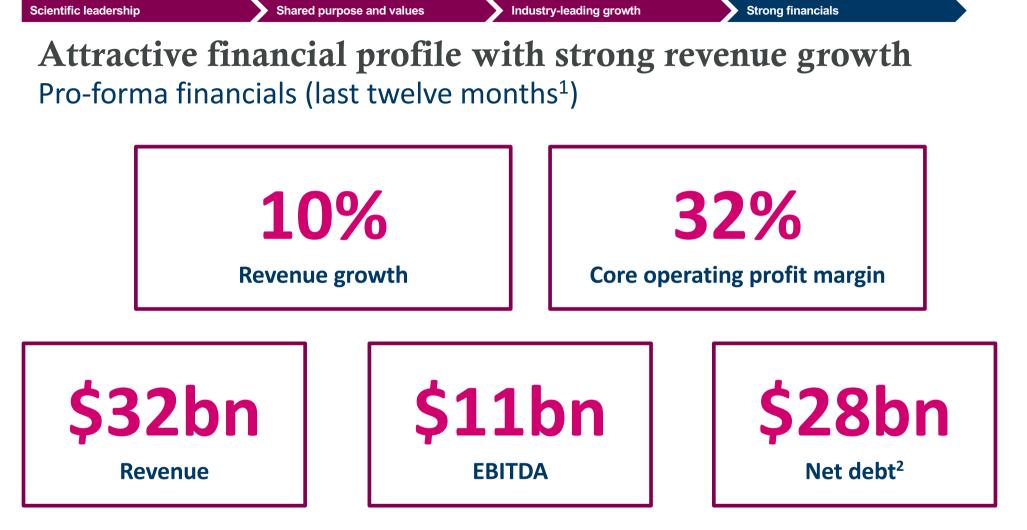
Industry-leading growth

Strong financials



Scientific leadership





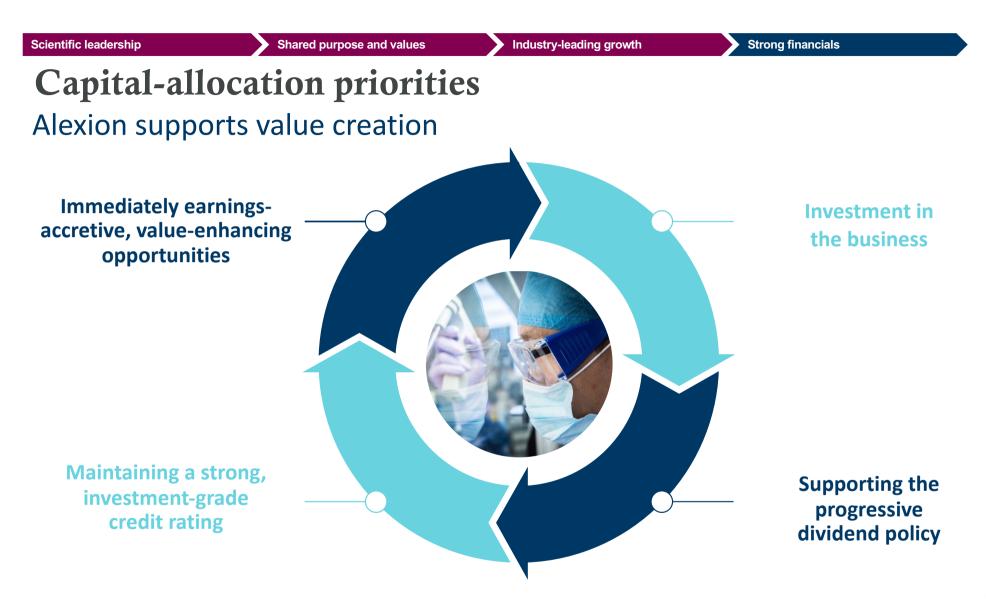
Source: Pro-forma financial metrics based on data from company financial statements over the last four quarters. EBITDA based on reported (AstraZeneca) and GAAP operating profit plus D&A (Alexion). All numbers are before any acquisition costs and synergies.

Notes 1: based on reported information as of 30/09/2020. Based on actual exchanges rates 2. includes c.\$13.5bn acquisition debt financing associated with cash consideration

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These statements are based on US GAAP financial results for Alexion These statements may be subject to amendment by AstraZeneca when stated in the Class 1 Circular and based on AstraZeneca's accounting policies."





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Details and timeline Anticipated closing in Q3 2021

Proposed key terms

- Price \$175 per share of Alexion
- Each shareholder of Alexion will receive
 - \$60 in cash
 - 2.1243 AstraZeneca American depositary shares (ADS) (Nasdaq: AZN)
- Total equity offer consideration \$39.4bn¹
 - \$13.5bn in cash, initially financed by underwritten bridge financing facility
 - \$25.9bn in AstraZeneca ADSs/shares

Total consideration \$39.4bn **Anticipated timeline**

- Following anticipated closing in Q3 2021, Alexion shareholders will own ~15% of the combined company
- Transaction subject to customary approvals, including by both sets of shareholders anticipated in Q2 2021
- Until closing, both companies will continue to operate as separate entities

Anticipated closing in Q3 2021





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Strong financials

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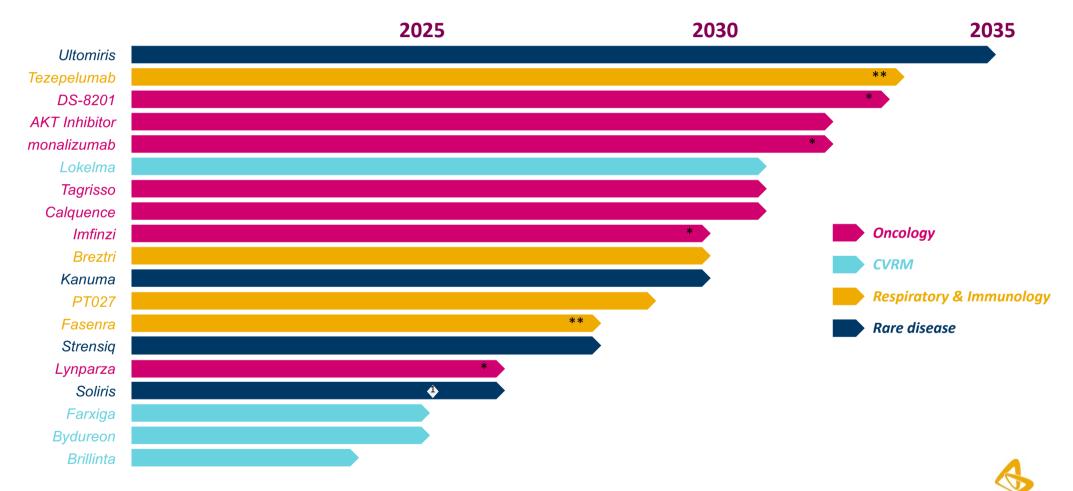
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- > Double-digit Core EPS accretion for the first three years
- > Strong cash flow, rapid debt deleveraging with an ambition to increase the dividend
- > Strong, investment-grade credit rating to provide strategic and financial flexibility



Growth sustainability underpinned by longevity beyond 2030



²⁶ Notes: US patent expiry. * Exclusivity period projected to extend beyond patent expiry due to patent restoration; ** Data exclusivity exceeds patent expiry. 1. Amgen IPR settled to grant Amgen a non-exclusive, royalty-free license to sell an eculizumab product in the US from March 1, 2025.