### VITA 34 – THE CELL BANK



### HOW TO BECOME MORE THAN A CELL BANK



## NEW ENHANCED MANAGEMENT BOARD - STRONGER FOCUS ON SALES & MARKETING





#### **Andreas Schafhirt (CFO)**

- Interim CFO, in the company since 2021
- 20+ years experience as CFO at listed and non-listed companies
- Responsible for Finance and Controlling

#### Jakub Baran (CEO)

- Co-founder, in the company since 2005
- 10+ years experience in Sales & IT at Blue Chip IT companies
- Responsible for Strategy, Operations, M&A, IR and Business Development

#### Tomasz Baran (CCO)

- In the company since 2010
- 10+ years experience at Big Pharma
- Responsible for Sales, Marketing, PR and Commercialization of Cell & Gene Therapies

### **EXECUTIVE SUMMARY – FULL YEAR 2021 RECAP**



Significant decline in earnings despite operational success in 2021

 Group Revenues
 EUR 28.4 m (+ 44.5 %)

 EBITDA
 EUR 0.8 m (- 83.5 %)

 EBIT
 EUR -3.1 m (- 255.7 %)

 Group Result
 EUR -3.9 m (- 372.8 %)

- Significant special items of EUR 5.1 m; Investments in new business areas EUR -1.3 m
- Core business with clearly positive EBITDA contribution of EUR + 7.3 m
- 4th quarter (peak of Omicron wave) with slowdown in new customer business...
- ...but continued strong development of recurring revenues (subscriptions): EUR 16.9m (2021 pro-forma)
- 2022 as transition year: focus on integration
- **Guidance 2022** Revenues EUR 68 75 m

EBITDA EUR -2 - +1 m

### **KEY FIGURES FOR THE BUSINESS DEVELOPMENT**



IFRS, in EUR '000	2021	2020	Δ
Revenues	28,419	19,668	44.5%
Gross profit on revenues	12,438	11,256	10.5%
EBITDA	814	4,936	-83.5%
EBITDA margin [%]	2.9%	25.1%	
EBIT	- 3,071	1,973	-255.7%
Result for the period	- 3,926	1,439	-372.8%
Earnings per share [in EUR]	- 0.63	0.35	-278.1%
Operating cash flow	2.727	3,531	-22.8%
Cash & cash equivalents (vs. 31 Dec. 2020)	33,298	10,396	220.3%

Note: Consolidation of PBKM FamiCord effective 8 November 2021

# ANALYSIS OF SPECIAL ITEMS: BUSINESS COMBINATION VITA 34/PBKM FAMICORD



- Business Combination of the two leading European Cell Banks
- Highly complex cross-border transaction results in significant transaction cost of EUR 2.7 m in total (Legal & General Advisory, Preparation & Implementation of Capital Increase, Exchange Offer and Squeeze-Out)
- Currently expected cost of integration in 2022: ~ EUR 0,5 m (net balance)

Expected Synergies in the next 3 years:2023: up to EUR 1 m

2024: up to EUR 2 m

2025: EUR 2.5 – 3 m

# ANALYSIS OF SPECIAL ITEMS: INVESTMENTS INTO NEW EXPANDING AREAS



"As by far the largest cell bank in Europe, we now have completely **new opportunities to shape our industry**, establish new products and services, and thus **position Vita 34 much more broadly and potently**.

We are more than certain that our investments today will establish the company in new, even far more attractive markets."

Jakub Baran, CEO Vita 34 AG

New expanding business areas: Cell & Gene Therapies (incl. CAR-T)

**CDMO** 

**Extensive investments already made** before Nov 2021: EUR 3 (5) m

✓ **Consolidated investments** in 2021: EUR 1.3 m

✓ **Planned investments** until 2024: EUR 30 m

# ANALYSIS OF SPECIAL ITEMS: HARMONIZATION OF ACCOUNTING ACC. TO IFRS 15



#### **IFRS 15, Revenue Recognition at Vita 34:**

- Special business model of Vita 34 comprises preparation on very long-term storage of samples
- Services either remunerated by long-term advance payments or subscriptions
- Special feature of PBKM contracts: certain termination rights during contract term
- Interpretation of IFRS 15 so far acc. to "Principle of Economic Effectiveness"
- Acc. To Vita 34 accounting guideline "Legal Principle" must be followed strictly:

Revenue Recognition for PBKM Subgroup had to be adjusted for past and current business!

Consolidated effect on Revenue and Earnings in 2021: EUR -1.6 m (cash neutral)

# **GUIDANCE TRANSITION YEAR 2022**



#### **GROUP REVENUES**

EUR 68 – 75 m

#### **GROUP EBITDA**

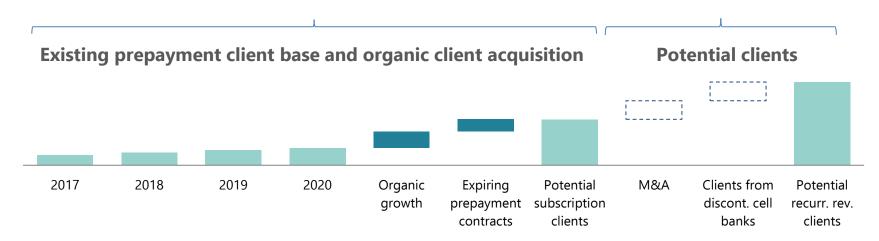
FUR -2 - +1 m

- Significant decline in earnings expected due to
   weakening of economic environment in relevant core markets
- Significant negative earnings impact from new expanding business areas to be expected
- Initial cost-cutting measures already initiated:
   mainly in areas of marketing, production, and from the post-merger integration process
- Potentially starting H2 2022: Recognition a greater volume of revenue from new contracts
   after appropriate adjustments to terms and conditions

# STRUCTURAL LEVERS: RECURRING REVENUES TO FUEL ORGANIC GROWTH



#### NUMBER OF CUSTOMERS IN SUBSCRIPTION PRICE MODELS



### STRUCTURAL LEVERS: NEW EXPANDING BUSINESS AREAS





### Cell & Gene Therapies (incl. CAR-T)

- 10 R&D Projects in progress
- ~4,200 patients in therapies
- 1st clinical trial of CARLA (CAR-T) approved
- 1st clinical trial of ALSTEM successful,
   2nd clinical trial in preparation

#### **CDMO**

- 3 synergistic R&D centers (DEU, POL & PRT)
- 15 own labs
   (Banking & ATMP manufacturing)
- Multiple CDMO contracts in EU & USA
   (e.g. NextCell, Dystrogen, Warsaw University)

New products and services will position Vita 34 much more broadly
> EUR 6 m invested so far, > EUR 30 m planned until 2024, incl. raise of EUR 20m share capital at
FamiCord Tx

### THE NEW VITA 34 AT A GLANCE



- ✓ No. 1 in Europe, No. 3 worldwide
- ✓ 850+ k biological samples stored
   + storage for ~300 k Ex-Cryo-Save clients
- √ ~95,2% of Group revenues\*

# Leading European cell bank



- √ ~227k recurring revenue clients (subscriptions)
- ✓ EUR 16,9 m of recurring cash revenues\* (subscriptions)
- ✓ Industry leading CLTV / CAC
- ✓ Steady recurring cash flow generation growing with double digit percentage

- ✓ Drug Manufacturing & Related
  - Experimental therapies
  - Multiple CMO contracts
  - Medical services
  - ~ 4,8% of Group revenues\*



### **Drug manufacturing & clinical trial portfolio**

- ✓ Clinical trials
  - Three own and three consortia projects
  - Ten clinical trials in five projects
  - Potential game changer for the valuation

### **GROUP FACILITIES – R&D CAPACITIES**



Location	Quality	Size		Location
Warsaw (POL)*	AABB, GMP	Very large	(>1,500 m <sup>2</sup> )	Sulzbach (GER)**
Cracow (POL)		Small	$(<150 \text{ m}^2)$	Poznan (POL)**
Cantanhede (PTG)*	AABB, GMP	Big	(250-500 m <sup>2</sup> )	New Jersey (USA)
Girona (ESP)	AABB, GMP	Mid	(150-250 m <sup>2</sup> )	Kiev (UKR)
Ankara (TUR)	GMP	Small	$(<150 \text{ m}^2)$	Nottingham (GBR)
Budapest (HUN)AABB		Mid	(150-250 m <sup>2</sup> )	Monthey (CHE)**
Bucarest (ROM)		Mid	(150-250 m <sup>2</sup> )	Stockholm (SWE)**
Cantone (CHE)	GMP Isolators	Small	$(<150 \text{ m}^2)$	Leipzig (GER)**
Leipzig (GER)	GMP FACT	Very large	$(>1,500 \text{ m}^2)$	Erlangen (GER)**
Madrid (ESP)	FACT	Mid	(150-250 m <sup>2</sup> )	+3 more**
Marly (CHE)	FACT	Small	$(<150 \text{ m}^2)$	
Istanbul (TUR)	GMP	Mid	(150-250 m <sup>2</sup> )	*ATMP Manufacturing
Ostrava (CZE)		Small	$(<150 \text{ m}^2)$	** storage only
London (GBR)		Mid	(150-250 m <sup>2</sup> )	
Rostock (GER)	GMP	Large	(500 - 1,500 m <sup>2</sup> )	

### **GROUP B2C/B2B SERVICES PORTFOLIO**



#### **Services offered (fully commercialized)**

**Technology applied (partially commercialized)** 

Cord blood banking

Placental blood banking

Cord tissue banking (segments, scraps, MSC isol./cult.)

Human semen banking

DNA banking

Adipose tissue banking (MSC / SVF)

Bone Marrow HSC separation & release (transplantation)

Peripheral blood HSC (transplantation)

Placenta (chorionic villi) MSC banking

Amniotic fluid MSC banking

Chimeras (myoblasts/MSC) manufacturing

#### **ATMP** manufacturing:

- ✓ MSC from Wharton Jelly
- ✓ MSC from adipose tissue

Placental MSC (chorion, decidua basalis, amnion)

Fibroblasts and other adherent cells

Dental pulp MSC

Bone marrow MSC

Exosomes from cord blood

Exosomes from adipose tissue

Viral vectors

**ATMP** manufacturing:

✓ CAR-T

### **GROUP R&D PROJECTS I**



Project	Description
Stroke Therapy	<ul> <li>✓ Bone Marrow CD34+ cells in ischemic stroke patients</li> <li>✓ Clinical trial planned Q12022</li> <li>Clinical trial Approved by National Ethics Committee for Clinical Research, waiting for National Authority of Medicines and Health Products approval</li> </ul>
Rescue Cord	<ul> <li>✓ Umbilical Cord Blood for newborns with Hypoxic Ischemic Encephalopathy</li> <li>✓ Hospital Exemption procedures planned for Q4 2021</li> <li>✓ Papers under construction</li> </ul>
MSCell Production	<ul> <li>✓ GMP grade manufacturing of MSC derived from adipose tissue and umbilical cord tissue for immunological disorders. General manufacturing approval for clinical trials</li> <li>✓ Manufacturing product authorization from National Authority of Medicines and Health Products received</li> </ul>
ARDS clinical trial	<ul> <li>✓ Umbilical cord-derived MSC for ARDS patients</li> <li>✓ Clinical trial in preparation, informed consent issues</li> </ul>

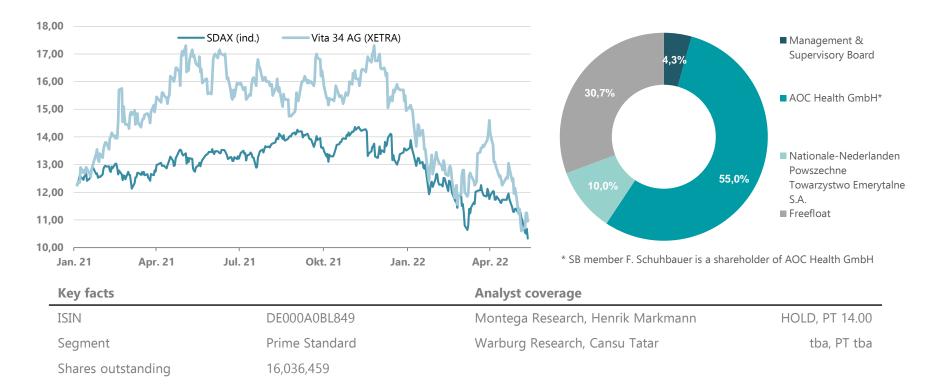
### **GROUP R&D PROJECTS II**



Project	Description
Circulate (Consortium)	<ul> <li>✓ UC MSC for cardiac diseases (Chronic Ischemic Heart Failure, Critical Limb Ischemia, Acute Myocardial Inf.)</li> <li>✓ Clinical trials phase II/III, 3 arms of total 315 patients</li> <li>✓ Clinical trials ended and concluded . Analysis of the results going on.</li> </ul>
ABC Therapy (Consortium)	<ul> <li>✓ MSC from adipose tissue for treatment of diabetic foot and in dermatology (Cutis laxa and scars)</li> <li>✓ Clinical trial phase II/IIIa ended Q3 2020 and pending HE procedures for diabetic foot</li> </ul>
BIOOPA (Consortium)	<ul> <li>✓ Bio dressing with UC MSC for treatment of epidermolysis bullosa and difficult scares</li> <li>✓ Clinical trial phase II/IIIa ongoing; R&amp;D currently ongoing</li> </ul>
ALSTEM	<ul> <li>✓ UC MSC for treatment of amyotrophic lateral sclerosis (ALS) and development of preclinical testing panel enabling better qualification of patients for cellular therapies</li> <li>✓ 1st clinical trial ongoing , 100% patients recruited , 2nd trial in preparation</li> </ul>
CD19_CAR-T	<ul> <li>✓ Purchased exclusive license from iCell Gene Therapeutics for use of CAR-T</li> <li>✓ technology in Europe, product development in PBKM, clinical trial in Warsaw Medical University</li> <li>✓ application for marketing authorization at EMA</li> </ul>

# SHARE PRICE DEVELOPMENT BURDENED BY TRANSITION YEAR







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