



GATT Hemostatic Platform

Next generation hemostats stopping complex surgical bleeding

May 2020 | GATT Technologies BV

Objectives of today's discussion

- Review hemostat and sealants markets
- Introduce GATT Technologies' platform and business
- Identify Business Development opportunities in Bay area

Executive Summary

Hemostats and sealants: attractive market, current products not suited to meet the need

- Hemostats and sealants address a surgical need to stop bleeding during surgery where mechanical measures are insufficient
- Substantial global market for hemostats and sealants (~\$3B globally), growing (~8% p.a), and continuing to grow as new therapeutic areas are developed
- All existing technologies present in the market have limitations there is no single solution yet
- Large market players have recognized existing technology limitations, and are investing in innovation and M&A

GATT has a game changing next generation platform addressing unmet need and expanding the market

- Developed activated polyoxazoline polymer synthetic technology platform, protected by strong IP portfolio, that generates superior performance at lower costs
- Technology platform generates ability to be a one stop shop, addressing all types of surgical procedures, and the full needs of surgeons
- GATT's first product, the GATT-Patch, is confirming its benefits compared to competition – faster time to hemostasis, better adhesion, lower COGs
- Surgeons in the US and EU are enthusiastic about GATT-Patch, and have confirmed the potential

GATT is executing on a two-year plan to CE (Q1 2022) and FDA (Q2 2023) approvals

- GATT has 2-year funding from J&JDC (ventures), Dutch Government OostNL and US PE fund, established experienced team, frozen GATT-Patch design, manufacturing and QMS capabilities
- Full (pre)-clinical plan is in place, and milestones for next 2 years have been set currently in execution mode
- First in human study expected in Q1 2021

GATT provides appealing investment opportunity

- GATT will likely seek additional funding post Q1 2022 to execute clinical studies required for FDA approval and ramp up to commercialization
- · Ongoing discussions with potential strategic partners



GATT has established knowledgeable and experienced management team



Geert van Gansewinkel, MSc, MBA

Chief Executive Officer

Successful life sciences executive

- Experienced entrepreneur and general manager
- 20+ years experience: Accenture, The Boston Consulting Group, Polaris, IQVIA
- Entrepreneurship experience: built Polaris EU/Asia business (software) participated in successful exit
- Experience in strategy, M&A, GM: country and region P&L responsibility, CEO GATT since March 2020



Johan Bender, PharmD

Founder and Chief Technology Officer

Serial entrepreneur and hemostasis expert

- 25 years R&D and general management experience
- Serial entrepreneur; built and sold multiple medical devices, pharma formulation IP, and med testing lab companies
- Founded GATT in 2011, developed GATT platform including patented technology
- CTO since March 2020



Vlad Hogenhuis, MD, MBA

Chief Medical Officer a.i Board member

Medical doctor and senior life sciences executive

- Former Chief Operating Officer Ultragenyx
- SVP and global franchise head GSK Specialty Care
- SVP and General Manager Merck, US and International markets



Rob Lips, MSc Chief Commercial Officer

Senior medical device product and business development executive

- 30 years of international management experience in medical device industry
- Former global sales and marketing for Philips x-ray systems and Philips Healthcare incubator
- Responsible for GATT commercial activities since 2013



Dr. Rosa Felix, PhD Chief Operational Officer

Scientific expert surgical sealants and hemostats

- PhD in development of polymer based surgical sealants and hemostats
- Experienced laboratory technician and researcher (Merck)
- Part of GATT team since 2012, responsible for supply chain (development, supply and production)



Backed by an experienced and senior non-executive board



Jaap Kampinga, MD Chairman

- CEO Immodulon Therapeutics
- CEO and founder of LaCaMedical
- Former chairman Xenikos, and Cellotec and R&D director of Quadrant Healthcare



Zeev Zehavi On behalf of investor J&J

- Vice President J&J, JJDC (investments)
- Investment director on the board on behalf of JJDC



Johan de Ruiter On behalf of investor OostNL

- Chairman Quanta Fluid solutions
- Initiator/founder of Fresenius Medical Care
- Founder of Auxilium GmbH



Sjaak Deckers On behalf of investor NGI

- · Former CEO GTX medical
- Founder, CEO and exit Sapiens Steering Brain Simulation
- Former R&D manager Philips Healthcare

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Market potential

GATT Platform

GATT business and progress

GATT investment opportunity

Surgeons need next generation products that are easier to use and more suitable in a broader set of applications

Surgeons on our advisory board tell us that there is a clear need for better, more efficient hemostatic agents:

- For **simple bleeding**, current standards of care such as pressure, cautery, and sutures are usually sufficient to stop the bleeding
- For **complex bleeding** and/or with **anticoagulated** patients, hemostats are often needed
- Existing hemostats are **limited** in functionality (long time to adhesion, limited adhesion, poor performance with anticoagulated blood), often require preparation time, and multiple products are needed in the operating room
 - Surgeons are looking for products that can cover a broader variety of existing and new surgery challenges
- Fast, effective hemostats allow the surgeon to continue operating and reduce blood loss and surgery time improving patient morbidity and mortality rates



~\$3B global market for hemostats and sealants, growing 8% annually

~\$3B market of surgical hemostats, sealants and glues, growing at 8% CAGR



Growth expected to continue

- Rising surgery volumes (particularly in the US)
- Technological advancements (MIS, robotic surgery, etc.)
- · Rising numbers of trauma cases and complex surgeries
- Growing prevalence of acute and chronic conditions (cardiovascular, obesity, cancer, etc.)
- · Shrinking blood supply/donor pool in many countries
- Increasing healthcare expenditure, drive for efficiency and quicker operating room turnover
- Expanding geriatric population

Source: MedMarket Diligence, LLC, company web sites, GATT research

NOTE: Market numbers being validated; in this analysis numbers adjusted downward, reports exist suggesting a total market size of \$5Bn



Existing products have substantial limitations

Category/generation	Launched in	Method of action	Limitation	Sample products
Gelatin/Collagen/ORC - blanks	~1970-1990	 Stimulate hemostasis by accelerating the formation of platelets, leading to fibrin clot 	 Long time to hemostasis - depends on blood factors to initiate and drive the clotting process Limited adhesion - can only control minimal bleeding Foam based products can lead to increased pressure Often (e.g. flowable matrices) require long (3-5 min) preparation time 	SurgicelGelfoamSurgifoam
Fibrin/thrombin – actives	~1990-2010	 Acting at later stage in the hemostasis cascade, producing more immediate hemostatic results 	 Tissue adhesion and time to hemostasis limited Limited effectiveness in anti-coagulated blood Relatively expensive (e.g. ~\$800-900 for fibrin sealant patch) 	Tisseel, EviclTachosil, EvarrestFloseal, Surgiflo
Synthetic (NHS-PEG) – actives, synthetic	~2015-2020	 Further driving ability to improve time to hemostasis, but also allowing to seal the bleeding tissue 	 Limitation in adhesion points, reducing tissue adhesion strengths, and limitsing the use in other applications (e.g. lung surgery) 	HemopatchVeriset



Existing market players have recognized the need to innovate

Market players are investing in related IP

>300 patents filed by corporates in the last 5 years...



Source: NLO, European patent and trademark attorneys

...as well as acquiring smaller players

Selected M&A deals

- 2016: Mallinckrodt \$175M purchase of Recothrom, Preveleak, Raplixa
- 2017: Terumo \$1.1B purchase of Angioseal, Femoseal, Vadosheet
- 2018: Stryker \$220M purchase of Hyperbranch, Adherus
- 2019 (canceled 2020): J&J \$400M purchase of Tachosil
- 2020 Baxter \$350 purchase of Seprafilm



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GATT has developed a game changing next generation technology platform protected by a strong IP portfolio

GATT has developed innovative and patented technology platform...

GATT NHS-POx (polyoxazoline) patented technology, enabling stronger wound adhesion compared to NHS-PEG (Veriset):



...protected by a strong IP portfolio



✓ covering full hemostat/sealant product scope
 ✓ freedom to operate: no blocking technology claims/ownership



GATT Platform has superior benefits

- Shorter time to hemostasis
- Short time to strong adhesion
- Able to handle stronger/larger, more complex bleedings
- Very stable: long in-use stability
- Versatile platform of multiple applications: powder for laparoscopic surgery and semipermeable for lungs
- Non-brittle, more pliable/flexible, allowing for broader application use
- Tunable functionality: variation in dosage possible, creating options for use
- Less swelling
- Low cost of goods

Preclinical tests confirm superior benefits at lower cost

Category	Metric	Tachosil (Fibrin based)	Veriset (Synthetic, NHS-PEG)	GATT-Patch (Synthetic, NHS-POx)	
Clinical efficacy	Time to hemostasis	3-6 minutes	<1 minute (75%)	20-60 seconds ¹	
	Adhesion strength	+	++	+++	
	Control of problematic bleeding, coagulated blood	+	++	+++	
	Irregular tissue adhesion	-	+/-	+++	
Safety	Time to absorb in body	12 weeks	4 weeks	4 weeks	
	Swelling control	++		++	
Ease of use	Preparation time (patch)	+++	+++	+++	
	Flexibility, pliable, "plug and play" ²	- (rigid form) + (if wet)	+/-	++	
	Laparoscopic use (Y/N)	Y	Y	Υ	
Cost	US Price	~\$600+	\$300-350 (only EU)	~\$200-250 ³	

1. 700 GATT-Patch prototypes tested and compared with competing products in the market. Products tested in 25 ex-vivo livers and more than 30 in-vivo pigs of which 8 in-vivo at US corporate strategic partners

2. Also known as "tuft-and-stuff"

3. \$200-250 is lowest offer price for GATT that makes sense with a healthy margin, given that GATT is much better Than Tachosil, we can offer at Tachosil's price level (\$600) or even go higher, and price for premium performance – allowing high margin



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GATT platform provides opportunity to be "one stop shop," covering all surgical procedures

GATT product pipeline

- GATT-Patch: fast hemostatic sealing patch for general (open) surgery
 Frozen Product Design
- GATT-Powder: sealing hemostatic <u>powder</u> for general surgery, laparoscopic use
 In development
- **GATT-Spray**: elastic sealant for general surgery and other applications (e.g. lung surgery)
- GATT-Tape: for preventing intestinal anastomotic leakage
- Other potential applications: bone regeneration, anti-adhesion, vascular, ophthalmology <u>R&D pipeline</u>

GATT platform: one stop shop



Surgeons are enthusiastic and confirm the potential of GATT

Surgeon statements after an *in-vivo* porcine study with the GATT-Patch:

"Fast, robust adhesion, I can instantly continue with surgery"

"Very flexible and pliable for irregular surfaces"

"I have been looking for a solution that is more pliable, flexible and can be used without preparation.."

"This [current] generation products works for simple bleedings. We don't have anything yet for **severe bleedings**, and this [GATT-Patch] does..."

"The patch is opening the door for a whole **new set of applications** like liver, lung, and spleen resections..."

"The powder application is going to be a game changer in laparoscopic surgery"

"This can be a platform of solutions **simplifying and speeding up** many different types of surgeries, improving patients' well being and saving lives"







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Key business milestones have been defined

	20	20	2021			2022				
	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
Finalization of pre-clinical/bio-compatibility testing										
Submission of CE and IDE documentation										
First in man (EU)										
CE study close out										
Submission CE documentation										
Next investment round (C-round)										
Start FDA trials										
CE mark										
FDA Approval									Jul	y 1 2023



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Several avenues exist for short- and long-term value creation

- Through series B round, GATT has secured financing to obtain CE approval end of Q1 2022
 - Parallel to CE approval, we are starting FDA approval preparations
- GATT might seek additional investment in the form of a C-series, to fund full execution of FDA and ramp up of commercial model
 - Considering several options with strategic buyers
 - Commercialization could be in the form of distribution model or with partner, considering several options
- · Several parties have shown interest for long term investment
 - Potential strategic buyers from large strategic partners in the US, EU, China and Japan
- Attractive deal value, current expectation value ~\$200-250M value based on comparable deals for one product
 - Significantly higher (\$300-400M) depending on platform options added

J&J, CR Bard, Baxter, Medtronic, Takeda
 Teijin, Fosun



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