



## Silver Ions in the Clinical Setting

Metal ions and metal ions in complexes or compounds have been used as disinfectant for centuries. Molecular mechanisms of silver ions on bacteria and viruses have been studied extensively<sup>1</sup>, multiple antimicrobial mechanisms proposed<sup>1,2</sup>, and antimicrobial efficacy found to increase with silver ion concentration3. Part of the ionic silver released precipitates with chloride or phosphate ions or becomes strongly bound in the form of inert complexes with albumins or macroglobulins. Metabolic pathways are similar irrespective of route and uptake4 once taken up into the blood. Silver is processed by the liver and excreted in the bile and later the faeces<sup>5</sup>.

Silver has been used in the clinical setting in a variety of applications - topical burn cream, wound dressings to treat burns and ulcers<sup>6</sup>, PMMA bone cement<sup>7,8</sup>, central venous catheters<sup>7</sup>, urinary catheters<sup>9</sup>, and dressings.

#### Agluna Surface Technology

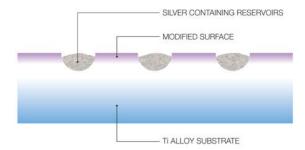
Agluna incorporates ancillary silver ions into the implant surface. Surface density is controlled by an ion exchange reaction. When the Agluna-treated implant comes into contact with body fluids, silver ions are released, helping to reduce the incidence of periprosthetic joint infection (PJI).

Agluna involves an electrochemical process using high voltage anodisation of the implant in phosphoric acid, forming an oxide layer integral with

<sup>1</sup> Thurman RB and Gerba CP, The molecular mechanisms of copper and silver ion disinfection of bacteria and viruses, CRC Critical Reviews in Environmental Control, 18(4), 295-315, 1989.

<sup>2</sup> Russell AD and Hugo WB, *Antimicrobial activity and action of silver*, Progress in Medicinal Chemistry, 31, 351-370, 1994.

the metal surface. Microscopic reservoirs are formed in the surface of the implant, containing hydrous titania into which silver ions are incorporated via a controlled ion exchange from a silver nitrate solution.



## Safety and Efficacy Evidence

Minimum Inhibitory Concentration (MIC): MIC of silver ions determined against clinical strains of Staph. aureus, Coagulase -ve Staph., Pseud. aerug. (gram -ve), E. coli (gram -ve), Enter. faecalus, Strep. pyogenes, Asper. fumigatus, Candida albicans and Klebsiella pneumonia<sup>10</sup>. MIC values consistent with those reported in literature. None of the tested clinical isolates exhibited results indicating resistance to silver.

#### • Antibacterial activity:

- ISO22196 study showed efficacy of Agluna articles against all nine organisms tested<sup>11</sup>.
- ASTM E2315 study provided quantitative evaluation of antimicrobial activity using timekill procedure. Antimicrobial effectiveness demonstrated against all microorganisms<sup>12</sup>.
- <u>Elution</u>: In vitro studies of Agluna-treated surface (up to five-year shelf-life period) showed silver ion elution kinetics consistent with benchmark CE-Marked Agluna METS device<sup>13</sup>.

<sup>&</sup>lt;sup>3</sup> Schierholz JM, Lucas LJ, Rump A and Pulverer, *Efficacy of silver-coated medical devices*, J. of Hosp. Infection, 40, 257-262, 1998.

<sup>&</sup>lt;sup>4</sup> Lansdown ABG, *A pharmacological and toxicological profile* of silver as an antimicrobial agent in medical devices, Adv Pharm Sci Volume doi: 10.1155/2010/910686, 2010.

<sup>&</sup>lt;sup>5</sup> Wan AT, Conyers R, et al, *Determination of silver in blood, urine, and tissues of volunteers and burn patients*, Clin Chem. 37(10), 1683-1687, 1991.

<sup>&</sup>lt;sup>6</sup> Burrell RE, *A scientific perspective on the use of topical silver preparations*, Ostomy Wound Management, Vol 49, 5A Suppl, 2003.

<sup>&</sup>lt;sup>7</sup> Spadaro JA, Webster DA and Becker RO, *Silver polymethyl methacrylate antibacterial bone cement*, Clin Orthop and Rel Research, 143, 887-892, 1979.

<sup>&</sup>lt;sup>8</sup> Sudmann E, Vik H, Rait M, Todnem K, Andersen KJ, Julsham K, Flesland O and Rungby J, *Systemic and local silver accumulation after total hip replacement using silver-impregnated bone cement*, Med Prog Tech, 20, 179-184, 1994.

<sup>&</sup>lt;sup>9</sup> Liedberg H and Lundeberg T, *Silver coating of urinary* catheters prevents adherence and growth of Pseudomonas aeruginosa, Urol Res., 17(6), 357-358, 1989.

Accentus Medical Reports ZRR\_ZB\_0005\_09, 2009;
ZRR\_ZB\_0007\_10, 2010; ZRR\_ZB\_0020\_09, 2009;
ZRR\_WA\_2658\_12, 2012.

<sup>&</sup>lt;sup>11</sup> Accentus Medical Report ZRR\_WA\_2965\_15, 2015.

<sup>&</sup>lt;sup>12</sup> Accentus Medical Report ZRR\_WA\_2790\_14, 2014.

<sup>&</sup>lt;sup>13</sup> Accentus Medical Report AGT07-DVer.059, 2015.

- Zone of Inhibition (ZoI): In vitro studies demonstrated ZoI around Agluna-treated coupons tested against MRSA<sup>14</sup>.
- Pharmacokinetics: In vivo study determined tissue distribution of eluted silver in organs/tissues over six-week time period. Cumulative silver concentration in all organs excluding liver and tibia was <2% of total silver delivered at all time points. Silver at the tested doses was not toxic to the study subjects<sup>15</sup>.
- Osseointegration: Literature research, in vitro and in vivo studies have evidenced early osseointegration of Agluna-treated implants<sup>16,17</sup>. Agluna gritblast surface showed no significant difference at 12 weeks relative to untreated gritblast surface. Histological investigations demonstrated normal osteoid tissue with no evidence of necrosis or osteolysis. No indication that Agluna surface adversely affects long-term osseointegration process of an uncemented titanium alloy implant.
- <u>Biocompatibility</u>: ISO10993-1 studies addressing a range of biocompatibility topics: cytotoxicity<sup>18</sup>; sensitization<sup>19</sup>; irritation<sup>20</sup>; acute systemic toxicity<sup>21</sup>; 1-, 4-, 12-, 26-week implant studies<sup>22</sup>; genotoxicity<sup>23</sup>; pyrogenicity<sup>24</sup>.

### **Agluna Manufacturing Process Validation**

Agluna process is operated by Accentus Medical at ISO13485:2016 accredited manufacturing facility. Processing plant has been validated in accordance with regulatory requirements. Performance Qualification study carried out demonstrating process validation with customer products.

# Clinical Experience

Agluna-treated limb salvage and complex revision devices successfully implanted since 2006: predominantly custom (patient specific) devices distributed in global markets by several specialist

device manufacturers operating in this sector. In 2011, Agluna-treated METS (Modular Endoprosthetic Tumour System) device (Stanmore Implants Worldwide Ltd, acquired by Stryker, Inc.) received a European CE mark.

In 2015, peer-reviewed clinical paper was published in Bone & Joint Journal<sup>25</sup>. This study reported a case-control study at Royal Orthopaedic Hospital (ROH), Birmingham, UK. 85 patients receiving Aglunatreated tumour implants (2006 to 2011) matched with 85 control patients receiving identical untreated implants (2001 to 2011). Key reported clinical outcomes were as follows:

- Reduction in post-operative infection rate from 22.4% (control) to 11.8% (Agluna);
- Success with DAIR (debridement, antibiotics and implant retention) increased from 31.6% (control) to 70% (Agluna);
- Persistent PJI necessitating device removal, amputation or chronic antibiotics suppression reduced from 15.3% (control) to 3.5% (Agluna);
- Success with two-stage revisions increased from 57.1% (control) to 85% (Agluna).

Post-operative clinical follow-up to 12 months, consistent with Agluna design objective.

A separate detailed evaluation of the ROH clinical data evidenced safety and performance of limb salvage and complex revision devices<sup>26</sup>.

Published studies conclude that silver-coated megaprostheses are safe and reduce the risk of PJIs in oncologic and septic revision surgery. Additionally they appear to offer a protective effect on implant survival in the event of infection, improving DAIR success rates and decreasing the need for salvage procedures such as amputations. These latter findings require validation in larger studies<sup>27</sup>.

In conclusion, potential risks related to Aglunatreated implants should be considered acceptable when weighed against the intended benefits for the patient.

<sup>&</sup>lt;sup>14</sup> Accentus Medical Report ZB-TEA-002, M22420, 2008.

<sup>&</sup>lt;sup>15</sup> Accentus Medical Report ZRR\_WA\_2782\_14, 2015.

<sup>&</sup>lt;sup>16</sup> Accentus Medical Reports ZTR\_WA\_0237\_14, 2014 (Coauthorship includes J. Parvizi); ZRR\_WA\_2744\_13, 2015.

<sup>&</sup>lt;sup>17</sup> ORS Poster, UCL / Accentus Medical, 2012.

<sup>&</sup>lt;sup>18</sup> Accentus Medical Report #189292 (WuxiApptec), 2013.

<sup>&</sup>lt;sup>19</sup> Accentus Medical Report #189287 (WuxiApptec), 2013.

<sup>&</sup>lt;sup>20</sup> Accentus Medical Report #189284 (WuxiApptec), 2013.

<sup>&</sup>lt;sup>21</sup> Accentus Medical Report #189286 (WuxiApptec), 2013.

<sup>&</sup>lt;sup>22</sup> Accentus Medical Reports #180054, #180055, #188188, #188187 (WuxiApptec), 2014.

<sup>&</sup>lt;sup>23</sup> Accentus Medical Reports #189283, #189285, #189288 (WuxiApptec), 2013.

<sup>&</sup>lt;sup>24</sup> Accentus Medical Report #189289 (WuxiApptec), 2013.

<sup>&</sup>lt;sup>25</sup> Wafa H, Grimer RJ, Reddy K, Jeys L, Abudu A, Carter SR and Tilman RM, *Retrospective evaluation of the incidence of early periprosthetic infection with silver-treated endoprostheses in high-risk patients*, Bone & Joint Journal, 97-B, 252-257, 2015.

<sup>&</sup>lt;sup>26</sup> Accentus Medical Report CSR\_CMU2011\_13H, 2014.

<sup>&</sup>lt;sup>27</sup> Sanz-Ruiz et al., Paper G53 to ICM2025.