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DOI
10.3389/fmats.2021.628633

Publication date
2021

Document Version
Final published version

Published in
Frontiers in Materials

Citation (APA)
Jahr, H., Li, Y., Zhou, J., Zadpoor, A. A., & Schröder, AK-U. (2021). Additively Manufactured Absorbable Porous Metal Implants: Processing, Alloying and Corrosion Behavior. Frontiers in Materials, 8, [628633]. https://doi.org/10.3389/fmats.2021.628633

Important note
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Additively Manufactured Absorbable Porous Metal Implants – Processing, Alloying and Corrosion Behavior

Holger Jahr¹,²*, Yageng Li³, Jie Zhou², Amir A. Zadpoor² and Kai-Uwe Schröder⁴

¹ Department of Orthopedic Surgery, Maastricht UMC+, Maastricht, Netherlands, ² Department of Biomechanical Engineering, Delft University of Technology, Delft, Netherlands, ³ Beijing Advanced Innovation Center for Materials Genome Engineering, School of Materials Science and Engineering, University of Science and Technology Beijing, Beijing, China, ⁴ Institute of Structural Mechanics and Lightweight Design, RWTH Aachen University, Aachen, Germany

Treating large bone defects is still a clinical challenge without perfect solution, mainly due to the unavailability of suitable bone implants. Additively manufactured (AM) absorbable porous metals provide unparalleled opportunities to realize the challenging requirements for bone-mimetic implants. Firstly, multi-scale geometries of such implants can be customized to mimic the micro-architecture and mechanical properties of human bone. The interconnected porous structure additionally increases the surface area to facilitate adhesion and proliferation of bone cells. Finally, their absorption properties are tunable to maintain the structural integrity of the implant throughout the bone healing process, ensuring sufficient loadbearing when needed and full disintegration after their job is done. Such a combination of properties paves the way for complete bone regeneration and remodeling. It is important to thoroughly characterize the biodegradation behavior, mechanical properties, and bone regeneration ability when developing ideal porous absorbable metal implants. We review the state-of-the-art of absorbable porous metals manufactured by selective laser melting (SLM), with a focus on geometrical design, material type, processing, and post-treatment. The impact of the latter aspects on absorption behavior, resulting mechanical properties, and cytocompatibility will also be briefly discussed. In comparison to their solid inert counterparts, AM absorbable porous metals (APMs) have shown many unique properties and hold tremendous potential to further optimize their application-specific performance due to their flexible geometrical design. We further highlight challenges in adopting AM APMs for future Orthopedic solutions.

Keywords: selective laser melting, absorbable implants, structure, corrosion, biomechanics

INTRODUCTION

Despite their self-healing abilities, large bone defects do not heal spontaneously and globally require millions of bone grafting procedures annually (Wu et al., 2014; Zhang L. et al., 2019). At present, 15–20% of the Western World population is 65 years or older and our aging society will soon face a dramatic increase in the demand for bone implants. This will cause increased
medical care costs as well as patient morbidity and mortality (Geetha et al., 2009; Fayaz et al., 2011; Loi et al., 2016). Current clinically applied bone grafts comprise autografts (a person’s own bone tissue), allografts (donor bone), and xenografts (non-human tissue). While devitalized allografts provide osteoconductive scaffolds of compromised mechanical stability for smaller defects (Giedraitis et al., 2014; Allsopp et al., 2016). Allografts and xenografts hold risks such as transplant rejection due to alloimmunity, transmitting diseases, infections, and compromised osteogenesis (Dimitriou et al., 2011). Therefore, autografts are the superior gold standard. However, they suffer from a limited supply, donor site morbidities, and complication rates of up to 40% (Van der Stok et al., 2011; Wang and Yeung, 2017).

Implants to treat critically-sized bone defects are, thus, particularly sought after on a bone grafts and substitutes market being valued at $4.15 billion by 2026 (Zhao et al., 2017; Polaris Market Research, 2020). Apart from being biocompatible, an ideal bone substitute might have a fully interconnected porous structure to allow for bone growth and possess bone-mimetic mechanical properties to provide sufficient support while avoiding stress shielding (Wen et al., 2001; Oh et al., 2003; Zadpoor, 2015). Highly porous metal implants were, therefore, introduced to improve biomaterial properties of traditional solid metals. With appropriately designed porosities, surface coefficients, and elastic modulus (Young’s modulus), they already mimic characteristics of cancellous bone (Amin Yavari et al., 2013; Matassi et al., 2013). Further advantages of 3D-printed biomedical metals over traditional implants include cost-efficiency, personalized design, and implantation site-specific tunable mechanical performance (Yan et al., 2018). However, like traditional solid permanent implants, they are still at risk of implant-associated infections or may require revision surgeries, with potential complications and unnecessary hospitalization and rehabilitation costs (Hexter et al., 2018).

Geometrically ordered AM porous metal implants with proper absorption profiles now offer these unique material properties and the possibility to fully regenerate bony defects with native bone. Absorbable implants would further allow regenerating bone tissue to replace the implant instead of growing around it or just into its interconnected pores. Given the implant’s complete disintegration, any risk of long-term infection would disappear with it – as opposed to permanently retained implants (Chen et al., 2014). So far, it has been challenging to produce porous absorbable biomaterials fulfilling all these requirements and the quest for ideal bone substituting materials is still booming (Oryan et al., 2014). We will specifically focus on absorbable porous implants manufactured by SLM technology, as AM biodegradable metals were recently reviewed (Li et al., 2020a), and summarize the impact of topological design- and material type-dependent differences in the corrosion behavior and corresponding mechanical properties. Furthermore, present shortcomings in adopting these implants for orthopedic applications and future opportunities will be briefly discussed.

### ADDITIVE MANUFACTURING OF ABSORBABLE METAL IMPLANTS

#### Additive Manufacturing Technologies

Current bone-substituting porous biomaterials may comprise polymers, ceramics, and metals (Do et al., 2015). Polymer-based biomaterials offer biofunctionalization and tailored biodegradation through design flexibility (Liu and Ma, 2004), while ceramic-based biomaterials are recognized for their favorable biodegradability and superior osteoconductivity (Seitz et al., 2005). For fully load-bearing applications in humans the former are too soft, while the latter are too brittle (Zhang L. et al., 2019). Metallic implants, on the other hand, have remarkable strength and energy absorption capacity, making them most suitable for such applications (Chen and Thouas, 2015). Complex design considerations are required to fabricate ideal porous metal bone implants. There are three main types of metal AM techniques, (i) directed energy deposition (DED), (ii) powder bed fusion (PBF), and (iii) binder jetting, which are described elsewhere in more detail (DebRoy et al., 2018; Li et al., 2020a). Generally, DED and PBF are both direct AM metal printing techniques, which can be further sub-divided by their heat source. Selective laser melting (SLM, Figure 1) is a laser-based PBF technology, also known as direct metal laser melting (DMLM) or laser powder bed fusion (LPBF). A more comprehensive overview on AM techniques for bio-inert metals also addresses their limitations (Li et al., 2020a). At present, LPBF is considered the most appropriate method for building complex porous structures and yet only absorbable porous metals meet all requirements to become ideal bone substitutes (Liu et al., 2017; Li et al., 2020a).

In contrast to bio-inert metals, AM absorbable metals (especially Mg and Zn) have low boiling temperatures and high chemical activities, posing new challenges for LPBF processing (Qin et al., 2019). Inappropriate processing conditions may cause defects, such as voids, lack of fusion, rough surfaces, severe residual stresses, and distortions. Even for Fe, evaporation can occur at high laser intensities (Kruth et al., 2004) and precise process control is, therefore, essential to successfully fabricate topologically ordered porous implants from absorbable metals by AM.

The type of PBF process applied potentially differentially affects the mechanical properties of porous metal products. Internal pores, inclusions, and cracks inside the struts of an implant can deteriorate its mechanical properties (Qin et al., 2019; Li et al., 2020a). Manufacturing defects of AM porous Zn may act as crack initiation sites, potentially shortening the fatigue life of an implant (Ahmadi et al., 2018). Optimizing the energy density tends to improve densification, and, thus, mechanical properties of SLM Mg alloys [e.g., AZ91 (Wei et al., 2014)], while different SLM process parameters affect the size and orientation of grains, thereby affecting mechanical properties as well (Manakari et al., 2016; Qin et al., 2020). Absorbable porous metals generally tend to have much finer grain sizes than their conventionally produced counterparts. This can improve
differ. Most Orthopedic and cardiovascular applications require depending on their anatomic location, healing rates of bone may be appreciated. Absorption of AM porous absorbable metals (PAM) are supposed to change with time. The projected different and constant mechanical properties, those of absorbable implants are supposed to change with time. The projected different and constantly increasing porosity. This suggests the lowest and highest degradation rates for Fe and Mg, respectively, with Zn having an intermediate rate. Biodegradation changes mechanical properties and fatigue behavior of AM PAMs. It decreases the fatigue life of SLM porous Mg and Fe (Christen et al., 2013; Nune et al., 2016): SLM porous absorbable Mg and Fe showed decreased yield strength and stiffness within 28 days of in vitro degradation under quasi-physiological conditions (i.e., in revised simulated body fluid, r-SBF). Currently, uncoated AM porous Mg and its alloys appear to corrode too fast and may prematurely fail in vivo.

Solid Zn implants corrode too slowly for most orthopedic applications and first 3D-printed Zn alloys recently improved this (Wen et al., 2018a,b). Surprisingly, after 28 days of biodegradation under the same conditions as for Mg and Fe, SLM porous Zn revealed increased mechanical properties as compared to as-built controls (Li et al., 2018a,b, 2020c),

Absorbable Metal Families
Recently, absorbable (ASTM F3160-16), or “biodegradable,” metals received increasing interest as they possess excellent mechanical properties and degrade in the human body without concerning long-term fatigue damage and revision surgery as compared to permanent counterparts (Do et al., 2015; Liu et al., 2017; Hermawan, 2018). The currently considered absorbable metal families comprise iron (Fe), magnesium (Mg), and zinc (Zn) (Li et al., 2020a; Wang et al., 2020). A comparison of the different mechanical properties of selected biomaterials to those of native bone illustrates the attractiveness of Mg, and its alloys, for load-bearing Orthopedic applications, and why Fe would benefit relatively the most from a decreasing Young's modulus through increasing porosity (Figure 1B).

In contrast to traditional permanent implants with almost constant mechanical properties, those of absorbable implants are supposed to change with time. The projected different and increasingly load-bearing phases of bone tissue regeneration have to keep pace with the corrosion speed-dependent decreasing mechanical support by the implant (Figure 1C).

Degradation behavior of AM absorbable bulk and porous metals are compared in Table 1. Importantly, the influence of loading on their corrosion behavior is currently poorly appreciated. Absorption of AM porous absorbable metals (PAM) can, however, be controlled not only by their material type or chemical composition (i.e., alloying), but also through their microstructure and geometrical design (Li et al., 2020a).

Corrosion Behavior of Absorbable Metals
Depending on their anatomic location, healing rates of bone may differ. Most Orthopedic and cardiovascular applications require mechanical support of at least three and 4 months, respectively, ideally being fully absorbed within 2 years (Zheng et al., 2014; Francis et al., 2015; Venezuela and Dargusch, 2019). Bone fixation devices are further suggested to have degradation rates of less than 0.5 mm/year (Chen et al., 2014; Zheng et al., 2014). In contrast, there is no widely accepted criterion for the corrosion rate of absorbable implants. Controlling degradation speed in situ is thus utmost important to avoid premature implant failure and proper corrosion testing of absorbable metallic materials in vitro is crucial. Logically, ideal degradation rates would match the location-specific bone regeneration pace (DebRoy et al., 2018). Therefore, means to control the corrosion rates of PAMs are highly sought after. Absorbable metals, and their corrosion products, must further be biocompatible, which will be briefly discussed in section “Biocompatibility and Clinical Application Potential”.

The electrode potential values of pure Fe, Mg, and Zn are −0.44, −2.37, and −0.76 V, respectively (Zheng et al., 2014). This suggests the lowest and highest degradation rates for Fe and Mg, respectively, with Zn having an intermediate rate. Biodegradation changes mechanical properties and fatigue behavior of AM PAMs. It decreases the fatigue life of SLM porous Mg and Fe (Christen et al., 2013; Nune et al., 2016): SLM porous absorbable Mg and Fe showed decreased yield strength and stiffness within 28 days of in vitro degradation under quasi-physiological conditions (i.e., in revised simulated body fluid, r-SBF). Currently, uncoated AM porous Mg and its alloys appear to corrode too fast and may prematurely fail in vivo.

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which was explained by formation of degradation products. Those are 5-times harder than AM Zn itself (Li et al., 2018a) and tightly bind to AM Zn (Li et al., 2020b). Vascular applications of zinc-based alloys were recently critically reviewed (Mostaed et al., 2018).

**Alloying-Dependent Corrosion Behavior**

During 28 days of *in vitro* corrosion, AM WE43 implants lost almost 20% of their weight, whereas AM Fe and Zn exhibited only 3% and 8% weight loss, respectively (Li et al., 2018b; Figure 1A). As Mg generally degrades too fast and Fe too

| Metal | Alloy | Unit cell design | Porosity (%) | Testing method | Elastic modulus (GPa) | Ultimate strength (MPa) | Yield strength (MPa) | References |
|-------|-------|-----------------|--------------|----------------|-----------------------|------------------------|---------------------|------------|
| Mg    | pure  | indentation     | 27–33        |                |                       |                        |                     | Ng et al., 2011a |
| Mg    | pure  | indentation     | 33–35        |                |                       |                        |                     | Ng et al., 2011b |
| AZ91D |       | tensile         | 296          | 254            |                       |                        |                     | Wei et al., 2014 |
| WE43  |       | tensile         | 45.7 ± 1.5   | 308            | 296.3 ± 2.5           |                        |                     | Zumdick et al., 2019 |
| Mg    |       | compression     | 51           |                |                       |                        |                     | Zhou et al., 2016 |
| Mg-1Zn|       | tensile         | 148          |                |                       |                        |                     | Wei et al., 2019  |
| Mg-2Zn|       |                | 71           |                |                       |                        |                     |            |
| Mg-4Zn|       |                | 60           |                |                       |                        |                     |            |
| Mg-6Zn|       |                | 45           |                |                       |                        |                     |            |
| Mg-8Zn|       |                | 44           |                |                       |                        |                     |            |
| Mg-10Zn|      |                | 63           |                |                       |                        |                     |            |
| Mg-12Zn|      |                | 74           |                |                       |                        |                     |            |
| Fe    | pure  | tensile         | 215.8 ± 20   | 411.5 ± 25     | 305.3 ± 20            | Song et al., 2014a    |                     |            |
| Fe    | pure  | tensile         | 208 ± 16     | 357 ± 22       | 256 ± 17              | Song et al., 2014b    |                     |            |
| Fe    | Pure  | compression     | 200          |                |                       |                        |                     | Montani et al., 2017 |
| Zn    | Pure  | tensile         | 14–32        | 132–138        | 108–122               | Wen et al., 2018b     |                     |            |
| Zn    | pure  | tensile         | 20.47 ± 5.71 | 137.9 ± 2.48   | 122.13 ± 2.61         | Wen et al., 2018a     |                     |            |
| Zn    | pure  | compression     | 117–133      | 93–110         | 79–90                 | Montani et al., 2017  |                     |            |
| Zn    | pure  | compression     | 146          |                |                       | Shuai et al., 2018c   |                     |            |
| Zn    |       | tensile         | 12.2 ± 2.4   | 61.3 ± 5.0     | 43.2 ± 3.1            | Yang Y. et al., 2018  |                     |            |
| Zn-1Mg|       | tensile         | 18.8         | 126.3 ± 3.6    | 74.1 ± 3.8            |                       |                     |            |
| Zn-2Mg|       |                | 25.2         | 161.8 ± 5.6    | 117.4 ± 5.4           |                       |                     |            |
| Zn-3Mg|       |                | 48.2 ± 4.2   | 222.3 ± 8.2    | 152.4 ± 4.8           |                       |                     |            |
| Zn-4Mg|       |                | 57.5 ± 4.8   | 166.4 ± 7.4    | 131.6 ± 7.5           |                       |                     |            |
| Zn-2Al|       |               | 141.7–192.2  | 121.4–170.5    |                       |                       |                     |            |
| porous Mg | WE43 | lattice  | 76.20        | compression    | 15                    | Kopp et al., 2019      |                     |            |
| porous Mg | WE43 | bending | 20–23        | compression    | 27                    | Witte et al., 2016    |                     |            |
| porous Mg | WE43 | diamond | 67           | compression    | 23                    | Li et al., 2018b       |                     |            |
| porous Fe | Fe-25Mn | cubic | 66.72 ± 2.3  | compression | 304.0 ± 7.4 | 89.2 ± 1.9 | Carluccio et al., 2020 |
| porous Fe | Fe-35Mn | sheet | 33.5 ± 1.7   | compression | 11.4                 | Li et al., 2019a       |                     |            |
| porous Fe | Fe    | diamond  | 84.6 ± 0.4   | compression | 221.7 ± 10.9 | 137 ± 8.4 |                          |
| porous Zn | Zn    | diamond  | 72.6 ± 2.3   | compression | 399.765              | Li et al., 2020c       |                     |            |

AZ91D (8.3–9.7% Al, ≤0.03% Cu, ≤0.005% Fe, ≤0.002% Ni, 0.35–1.0% Zn, 0.15–0.5% Mn, ≤0.1% Si, 0.02% other metals); WE43 (UNS M18430; rare earths 2.4–4.4%, 3.7–4.2% Y, 0.4% Zr).
| Material | Alloy composition | Unit cell design | Porosity (%) | Pore size (mm) | Duration (days) | Testing medium | Control | Weight loss (mm/year) | CRI (mA/cm²) | CRE (mm) | References |
|----------|-------------------|------------------|--------------|---------------|----------------|----------------|---------|----------------------|--------------|----------|------------|
| Mg       | Mg                |                  |              |               | 10             | SBF            |         | 13.30                |              |          | Zhou et al., 2016 |
|          | Mg-1Sn            |                  |              |               |                |                |         |                      |              |          |            |
|          | Mg-3Sn            |                  |              |               |                |                |         |                      |              |          |            |
|          | Mg-5Sn            |                  |              |               |                |                |         |                      |              |          |            |
|          | Mg-7Sn            |                  |              |               |                |                |         |                      |              |          |            |
| AZ61     |                   |                  | 6            |               |                | SBF            | 1.2-2.7 |                      |              |          | He et al., 2017    |
| ZX60     | 7                 | SBF              |              |               | 2.1            | 44.2           |         |                      |              |          | Shuai et al., 2018b |
|          | 0.2Cu             |                  |              |               | 2.4            | 60.4           |         |                      |              |          |            |
|          | 0.4Cu             |                  |              |               | 2.5            | 85.3           |         |                      |              |          |            |
|          | 0.6Cu             |                  |              |               | 17             | 485.7          |         |                      |              |          |            |
|          | 0.8Cu             |                  |              |               | 30             | 827.5          |         |                      |              |          |            |
| ZK30     | SBF              |                  |              |               |                |                |         | 131 ± 14             |              |          | Shuai et al., 2018a |
| ZK30-1Al |                  |                  |              |               |                |                |         |                      |              |          |            |
| ZK30-3Al |                  |                  |              |               |                |                |         |                      |              |          |            |
| ZK30-5Al |                  |                  |              |               |                |                |         |                      |              |          |            |
| ZK30-7Al |                  |                  |              |               |                |                |         |                      |              |          |            |
| Mg-6Zn-0.5Zr |            |                | 28           |               |                | SBF            | 1.58 ± 0.21 | 36.2 ± 2.3 |              |              |          | Yang Y. et al., 2018 |
| Fe       | Fe                | HANKS            |              |               |                |                |         | 6.2 ± 0.1            |              |          | Caruccio et al., 2019 |
|          | 2Pd-2.5bredigite  |                  | 21           |               |                | SBF            | 0.22      | 17.78                |              |          | Gao et al., 2019    |
|          | 2Pd-5bredigite    |                  | 21           |               |                | SBF            | 0.38      | 31.62                |              |          |            |
|          | 2Pd-10bredigite   |                  | 21           |               |                | SBF            | 0.5       | 39.81                |              |          |            |
|          | 4Pd-2.5bredigite  |                  | 21           |               |                | SBF            | 0.41      | 34.67                |              |          |            |
|          | 4Pd-5bredigite    |                  | 21           |               |                | SBF            | 0.6       | 50.12                |              |          |            |
|          | 4Pd-10bredigite   |                  | 21           |               |                | SBF            | 0.76      | 63.09                |              |          |            |

(Continued)
### TABLE 1B | Continued

| Material | Alloy composition | Unit cell design | Porosity (%) | Pore size (mm) | Duration (days) | Testing medium | Control | Weight loss (mm/year) | CRI (mA/cm²) | CRE (mA/cm²) | References |
|----------|------------------|-----------------|--------------|---------------|----------------|----------------|---------|-----------------------|-------------|-------------|------------|
| Zn       | Zn               | 76.20           | 1131         | 21            | HANKS          | CO₂            | 40.20   | 0.038–0.04           | 1.6–4.29    | Qin et al., 2020 |
| Zn       |                  |                 |              |               |                |                |         | 0.081                 | 7.76        | Shuai et al., 2018c |
| Zn-2Ag   |                  |                 |              |               |                | SBF            | 0.086   | 0.107                 | 1.47        |             |
| Zn-4Ag   |                  |                 |              |               |                | SBF            | 0.114   | 1.47                  |             |             |
| Zn-6Ag   |                  |                 |              |               |                | SBF            | 0.133   | 13.94                 |             |             |
| Zn-8Ag   |                  |                 |              |               |                | SBF            | 0.137   | 13.94                 |             |             |
| Zn-2Al   |                  |                 |              | 21            | SBF            |                | 0.13–0.16 | 7.07–11.75            |             | Shuai et al., 2019a |
| Zn-2Al   |                  |                 |              | 14            | SBF            |                | 0.14     | 5.86                  |             | Yang Y. et al., 2018 |
| Zn-4Al   |                  |                 |              | 28            | SBF            |                | 0.18 ± 0.03 | 9.24 ± 1.21          |             |             |
| Zn-6Al   |                  |                 |              | 28            | SBF            |                | 0.14 ± 0.01 | 5.86 ± 1.42          |             |             |
| Zn-8Al   |                  |                 |              | 28            | SBF            |                | 0.13 ± 0.03 | 4.63 ± 0.95          |             |             |
| Zn-2Mg   |                  |                 |              | 28            | SBF            |                | 0.10 ± 0.02 | 3.62 ± 0.76          |             |             |
| Zn-4Mg   |                  |                 |              | 28            | SBF            |                | 0.11 ± 0.04 | 3.71 ± 0.87          |             |             |

| porous Mg | WE43 | 76.20 | 1131 | 21 | DMEM | CO₂ | 40.20 | 0.23 | 21–61 | Kopp et al., 2019 |
| porous Mg | WE43 | 76.20 | 1131 | 21 | HANKS | bioreactor | 25 |       | Witte et al., 2016 |
| porous Mg | WE43 | 76.20 | 1131 | 21 | r-SBF | HEPES | 20.70 | 0.23 | 21–61 | Li et al., 2018b |

| porous Fe | Fe   | 66.72 | 400  | 28 | SBF |                | 5.3 | 0.09 ± 0.02 | 7.38 ± 3.21 | Shuai et al., 2020 |
| porous Fe | Fe-25Mn | 66.72 | 400  | 28 | SBF |                | 13.4 | 0.23 ± 0.05 | 51.25 ± 7.52 |             |
| porous Fe | Fe35Mn | 66.72 | 400  | 28 | SBF |                | 69.7 | 0.14   |             |             |
| porous Fe | Fe   | 66.72 | 400  | 28 | SBF |                | 70.3 | 0.17   |             |             |
| porous Fe | Fe   | 66.72 | 400  | 28 | SBF |                | 58.9 | 0.11   |             |             |
| porous Fe | Fe   | 66.72 | 400  | 28 | SBF |                | 73%  | 0.13   | 102.8 ± 19.2 | Li et al., 2018a |

| porous Zn | Zn   | 76.20 | 1131 | 21 | HANKS | bioreactor | 11.9 | 0.17 |             |             |
| porous Zn | Zn   | 76.20 | 1131 | 21 | SBF |              | 68.5 | 0.14 |             |             |
| porous Zn | Zn   | 76.20 | 1131 | 21 | SBF |              | 62.0 | 0.13 |             |             |
| porous Zn | Zn   | 76.20 | 1131 | 21 | SBF |              | 62.0 | 0.07 |             |             |

AZ61 (M16600; Mg 94%, Zn 4.8–6.2%, Zr 0.45%); AZ91D (8.3–9.7% Al, ≤0.03% Cu, ≤0.05% Fe, ≤0.002% Ni, 0.35–1.0% Zn, 0.15–0.5% Mn, ≤0.1 Si, 0.02% other metals); WE43 (UNS M18430; rare earths 2.4–4.4%, 3.7–4.2% Y, 0.4% Zr); ZK61 (Mg 93%, Al 6%, Zn 1%); DMEM, Dulbecco’s Modified Eagle’s Medium (NaHCO₃-buffered with sodium pyruvate and L-glutamine); HANKS, sodium bicarbonate-buffered salt solution (0.35 g/L NaHCO₃, 1 g/L glucose); SBF (142 mM Na⁺, 5 mM K⁺, 1.5 mM Mg²⁺, 2.5 mM Ca²⁺, 148.8 mM Cl⁻, 4.2 mM HCO₃⁻, 1 mM HPO₄²⁻, TRIS-buffered) (Kokubo and Takadama, 2006); r-SBF (HEPES-buffered SBF with 103 mM Cl⁻, 27 mM HCO₃⁻, and additional 0.5 mM SO₄²⁻) (Cyane et al., 2003). CRI, Corrosion rates determined gravimetrically from immersion tests; CRE, Corrosion rates determined electrochemically. Tests requirements: bone substitute, not known; bone fixature < 0.5 mm/year (Chen et al., 2014); stents < 0.02 mm/year (Bowen et al., 2013).
slowly, recent efforts were mainly directed toward tuning their corrosion behavior by alloying. This may lead to grain refinement to improve degradation resistance of Mg, but can generate second phase(s) or cause grain boundary segregation, both of which can accelerate galvanic degradation. Biodegradation rate of SLM Mg was reduced by alloying with 1% Sn, while higher percentages of tin increased corrosion rates likely due to second phase(s) overshadowing the grain refinement effects (Zhou et al., 2016). Similar results are reported for SLM ZK30 (Shuai et al., 2018a). However, alloying Mg with aluminum (Al), like in AZ91 (8.3–9.7% Al) or AZ31B (2.5–3.5% Al), reduces its corrosion rate at the expense of its biocompatibility (Ghosh et al., 2020). A comparison of degradation rates of Fe-, Zn-, and Mg-based alloys (including AZ31B) was recently published, illustrating the improved corrosion resistance of Mg-0.8Ca and Mg1Zn (Mei et al., 2020).

Corrosion rates of Fe-Mn alloys are reported to be higher than those of AM pure Fe (Li et al., 2019b; Shuai et al., 2020) because of increased galvanic degradation. Fe-20Mn was recently reported to have a decreased corrosion rate as compared to the pure metal, though (Mei et al., 2020). Addition of silver to Zn increased its corrosion rate, while, interestingly, addition of Mg to Zn either decreased or increased it, probably due to concentration-dependent different impacts of grain refinement and second phase (Shuai et al., 2018c; Yang Y. et al., 2018; Mei et al., 2020).

**Impact of the Composition of the Test Solution on Corrosion Behavior**

Published discrepancies between absorbable metal corrosion behavior likely result from non-comparable test conditions (Mei et al., 2020). To mimic human body physiology, it remains meaningful to use buffers to keep pH values constant (Törne et al., 2017). The addition of buffers like HEPES, however, can destabilize protective layers and accelerate Mg degradation by a magnitude. In absence of corrosion accelerating Tris–HCl, formation of uniform protective corrosion product layers slows corrosion rates in DMEM as compared to SBF (Liu et al., 2019). Using HEPES promoted the formation of passive layers in NaCl solution to reduce corrosion rates, while it showed opposite effects in SBF (Liu et al., 2019).

Of the few studies comparing the corrosion rates of these metals directly under similar conditions, a recent study analyzed all three metals in two simulated physiological solutions, and reported their exact composition for the 28-day immersion tests (Dong et al., 2021). In essence, corrosion rate was initially slower in Dulbecco’s Modified Eagle’s Medium (DMEM) for all three metals, with Mg showing the fastest burst corrosion rate in both SBF and DMEM. Intriguingly, the ranking of the relative corrosion speed changed over time, showing an increasing corrosion resistance of Mg after the initial 24 h, which confirms earlier data (Li et al., 2018b, 2020a). Typically, the corrosion rate of Mg is decreased as a function of increasing complexity of the medium, which indicates an urgent need to develop standard protocols for corrosion tests of absorbable metals (Mei et al., 2020). With a focus on Mg, recently a selection guide for corrosion media for absorbable metals, summarizing their advantages and shortcomings, was reported (Mei et al., 2020).

**Design- and Post Manufacturing-Dependent Corrosion Aspects**

Bone tissue is characterized by a gradual change in porosity from the outer cortical, compact bone toward an inner spongy tissue, with long bones being typical examples. This porosity and directionality is highly graded and dependent on local mechanical stimuli (e.g., strain energy density) (Campoli et al., 2012; Christen et al., 2013; Hazrati Marangalou et al., 2013; Zadpoor et al., 2013; Geraldes and Phillips, 2014; Nunes et al., 2016). It is tempting to mimic these natural structures in AM porous implants as they will eventually be surrounded by new bony tissue of similar micro-architecture and biomechanical characteristics (Nune et al., 2016; Zhang X.-Y. et al., 2019).

AM porous absorbable metals showed a strongly location-dependent corrosion behavior. The core of cylindrical AM porous Mg implants corroded much more prominently than their periphery (Li et al., 2018b); potentially due to limited diffusion upon accumulation of degradation products within the narrow spaces between struts, where Mg ions locally may cause crevice-like corrosion. In contrast, AM porous Zn corroded primarily localized in a contact zone with the reaction vial, likely due to hindered diffusion (Li et al., 2020c). Despite further identical designs and testing methods, in vitro degradation of AM porous Fe occurred more prominently on its periphery than in its core (Figures 1C; Li et al., 2018a). We recently used geometrical design to control biodegradation of SLM porous Fe to show that corrosion rates were directly proportional to its porosity. Furthermore, its biodegradation profiles could be tuned using functionally graded designs (Li et al., 2019a). This was later confirmed for Zn (Li et al., 2020c), underscoring the importance of rational design strategies for AM PAM implants (Table 1).

Similar to inert AM porous metals, yield strengths and elastic modulus of AM PAMs are directly related to their porosity through a power law (Li et al., 2019a, 2020c). Two types of unit cells are prevalent: (i) lattice structures and (ii) sheet-based structures (Figure 1C) such as minimal surface designs (Zadpoor, 2019a). The former can be subdivided into bending- and stretch-dominated structures; with bending-dominated types generally having a higher energy absorption ability, while the latter structures show higher stiffnesses and yield strengths (Deshpande et al., 2001; Zadpoor, 2019b). SLM porous Fe cubic unit cell-based designs possess higher yield strengths than diamond unit cell types (Li et al., 2018a; Shuai et al., 2020). AM Zn of constant porosity also showed unit cell-dependent mechanical properties (Lietal., 2020), but unit cell types also likely determine failure modes of scaffolds under compression (Mazur et al., 2016; Bobbert et al., 2017). SLM-based AM of functionally graded porous Fe and Zn of precisely controlled geometries were reviewed (Li et al., 2020a), realizing implants of trabecular bone-like mechanical properties.
Logically, HA coating on AM Fe scaffolds decreased its ion level of passivity on the metal surface (Ralston et al., 2010). The relationship between grain size and corrosion rate may depend on the Fe with larger grains (Li et al., 2018a), as the relationship of pure Zn, WE43, and pure Fe, using the same osteoblast-like cells and identical assays (Li et al., 2018a, 2019a, 2020d) to reveal a decreasing cytocompatibility in this order under these conditions. Like with corrosion testing, present cytocompatibility studies are hardly comparable. Limitations of most current studies further include very short in vitro evaluation periods and highly cell type-dependent outcomes (Li et al., 2020a).

In summary, SLM Fe appears to be relatively cytotoxic (Li et al., 2018a), potentially limiting its applications, while pure porous Mg corrodes fast (Hermawan, 2018). Currently, Zn and its alloys, seems to have an intermediate absorption rate and a reasonable cytocompatibility, which is currently based on only a handful of reports (Table 1), primarily in a cardiovascular context (Bowen et al., 2016; Drellich et al., 2017; Levy et al., 2017).

**Biocompatibility and Clinical Application Potential**

Significant amounts of the body's Mg and Zn content are stored in the inorganic bone matrix. The daily recommended intake of Mg is about 40–45 times higher than that of iron and zinc (Habibovic and Barralet, 2011; Florencio-Silva et al., 2015), suggesting its good tolerability (Figure 1B). Current clinical applications of magnesium-based implants in Orthopedics were recently reviewed comprehensively (Wang et al., 2020).

Several in vitro studies, as well as small animal studies, reported promising results on using iron as scaffolding material for cardiovascular and Orthopedic applications (Li et al., 2020a). Also a recent large animal model using iron-based degradable sponge-like implants manufactured by a powdery metallurgical replication method underscored its potential for self-degrading, high load-bearing bone replacements (Wegener et al., 2020). Smart designs can further increase corrosion rates of Fe toward avoiding long-term side effects (Rahim et al., 2018), but own studies on porous SLM iron implants prompt to caution (Li et al., 2018a, 2020a). Corroding iron fragments are further known to migrate through the body, causing chronic inflammation (Rahim et al., 2018).

Zinc alloys with favorable corrosion kinetics and biocompatible degradation products hold several promises for e.g., stent applications (Rahim et al., 2018).

 Appropriately modified magnesium-based implants are already marketed for cardiovascular applications (e.g., stents) and Orthopedic applications (e.g., compression or interference screws) and recent studies claimed they are resorbed to 95% within a year (Rahim et al., 2018). To better predict their translational potential, improved standards are not only needed for corrosion testing, but also for assessing their cytocompatibility in vitro. Recommendations to improve ISO 10993-5/-12 for Mg (Jung et al., 2019), and a recent technical corrigendum to ISO 10993-1:2009 (i.e., ISO 10993-1:2018) now containing additional information on the evaluation of, among others, absorbable materials, are important steps to facilitate the translational development of novel absorbable biomaterials.

A series of identically designed cylindrical porous SLM implants were used to systematically evaluate the cytotoxicity of pure Zn, WE43, and pure Fe, using the same osteoblast-like cells and identical assays (Li et al., 2018a, 2019a, 2020d) to reveal a decreasing cytocompatibility in this order under these conditions. Like with corrosion testing, present cytocompatibility studies are hardly comparable. Limitations of most current studies further include very short in vitro evaluation periods and highly cell type-dependent outcomes (Li et al., 2020a).

In summary, SLM Fe appears to be relatively cytotoxic (Li et al., 2018a), potentially limiting its applications, while pure porous Mg corrodes fast (Hermawan, 2018). Currently, Zn and its alloys, seems to have an intermediate absorption rate and a reasonable cytocompatibility, which is currently based on only a handful of reports (Table 1), primarily in a cardiovascular context (Bowen et al., 2016; Drellich et al., 2017; Levy et al., 2017).
Other recent review articles focused on the design and degradation (Shuai et al., 2019b; Li et al., 2020a) of absorbable Orthopedic implants, corrosion testing methods (Mei et al., 2020; Dong et al., 2021), biomaterial- stem cell interactions (Gao et al., 2017), the role of magnesium ions in bone repair (Wang et al., 2020), and the clinical translation of absorbable metals (Han et al., 2019; Wang et al., 2020).

In this mini review, we highlighted recent developments in SLM-based AM of PAM implants as they hold incredible potential to satisfy all basic functional requirements of an ideal bone-substituting implant. Their microstructural design holds a plethora of opportunities to fine-tune their corrosion behavior to specific clinical applications. To facilitate the transition from basic research into clinics, and to close remaining knowledge gaps in this rapidly evolving field, better standards for reporting corrosion test results and cytocompatibility data are needed though. Thus, a lot needs to be improved before widespread clinical application of such materials is feasible.

Due to lacking data, it is currently still largely unclear if in vitro corrosion rates of AM PAMs are by any means predictive of the in vivo situation. Next to reporting test solution details, it may be better to report weight loss percentages in future evaluations in addition to degradation rates as mm/year, because the latter unit makes more sense for bulk absorbable metals to which it is usually applied. For porous structures, the ratio of surface area to weight is much higher than for bulk counterparts and changes constantly during the degradation process. For bulk Mg, in vitro degradation rates are generally 1–4 times higher than values obtained in vivo, but are largely depending on the used alloy and in vitro testing conditions (Sanchez et al., 2015). To tune absorption rates of AM porous metals where needed, it is imperative to understand how in vitro corrosion is affected by chemical composition, microstructure, topological design, and surface conditions.

To improve mechanical properties, (i) Mg, Fe, and Zn alloys with higher mechanical properties need to be developed specifically for AM, (ii) as of yet underappreciated stretch-dominated or minimal surface designs as well as, and (iii) functionally graded structures combining different unit cell types and sizes should be considered. Transformation hardening may more likely work for Fe-based absorbable alloys than for Mg and Zn, or derivatives (Li et al., 2020a). As AM is a net-shape manufacturing technique, most strengthening mechanisms usually applied to bulk metals will not be applicable and accessibility inside 3D porous structures will be severely restricted. Rapid solidification during SLM results in finer grains, with size adjustment potential by layer-wise thermal history controlling. Process optimization and post-AM heat treatments may also improve mechanical properties. Alloying pure Fe with other elements might improve its biocompatibility, mechanical properties and biodegradation rate. Exploring Mg-based porous metallic glasses could result in AM Mg-based porous biomaterials with simultaneously improved corrosion resistance and mechanical properties. Zn-based mechanical properties may benefit from alloying Zn with other elements, particularly Mg and Ca. Better surface polishing methods and procedures are also needed (Li et al., 2020a). Creep and aging of AM Zn-based materials and their influences on the performance of these materials need to be further investigated. Different loading regimens (e.g., compression-tension, tension-tension, bending, and torsion) should be applied to all AM PAMs. Finally, future AM porous implants should not only consider initial bone-mimetic mechanical properties, but also anticipate dynamic changes during bone regeneration as well as parallel occurring deterioration of the implant properties. Thus, requiring much more sophisticated approaches.

This also holds for biocompatibility testing of novel implants, a detailed discussion of which is beyond our scope. However, good osteointegration would be a prerequisite to a successful clinical translation. As was shown for iron and zinc (Ray et al., 2018; Yang H. et al., 2018), novel coatings may be considered to potentially tune implant degradation and bone formation rates (Yu et al., 2017; Kang et al., 2019), too. It is envisioned that appropriate (functional) coatings will improve bone-forming abilities of all future AM PAMs. The few current in vivo studies unanimously reported new bone formation around SLM porous absorbable metal implants, but longer studies are desirable (Wang et al., 2020).

The “materialome” should be established to guide novel developments instead of pursuing a trial-and-error approach; (ii) the influence of the geometrical design on mechanical properties, corrosion behavior, cell-material-adhesion, and osseointegration should be systematically investigated by experimental and numerical approaches; (iii) the big gap between in vitro and in vivo results has to be closed. The plethora of educt and process parameters will become a serious challenge which may require real-time monitoring of the melt pool and powder bed in situ (Li et al., 2020a).

General recommendations for potentially interesting future work include (i) considering machine learning approaches to improve the selection of the process parameters and to better appreciate the complex relationships between chemical composition, lattice structure geometry, processing parameters, microstructure, and the resulting mechanical properties. A bright future awaits SLM porous absorbable metal implants when satisfying basic functional requirements of an ideal bone-substitute becomes achievable. Yet, more sophisticated round-robin evaluations are required to close current gaps in our knowledge.

**AUTHOR CONTRIBUTIONS**

HJ drafted, revised the manuscript, and submitted the final version. YL, JZ, AZ, and K-US contributed specific sections and critically co-revised the manuscript. All authors read the final version and agreed to its submission.

**FUNDING**

This work was supported by the Federal Ministry of Education and Research (BMBF) and the Ministry of Culture and Science.
of the State of North Rhine-Westphalia (MKW) under the Excellence Strategy of the Federal Government and the Länder (OPSF597). Financial support from the PRosPERoS project, funded by the Interreg VA Flanders–Netherlands Program (CCI grant no. 2014TC16RFC046), and the Medical Faculty (IZKF) of the RWTH Aachen University is also acknowledged.

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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