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DEGRADABLE ZINC MATERIAL CHARACTERISTICS AND ITS INFLUENCE ON
BIOCOMPATIBILITY IN AN IN-VIVO MURINE MODEL

By

Roger J. Guillory II

A DISSERTATION

Submitted in partial fulfillment of the requirements for the degree of

DOCTOR OF PHILOSOPHY

In Biomedical Engineering

MICHIGAN TECHNOLOGICAL UNIVERSITY

2019

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This dissertation has been approved in partial fulfillment of the requirements for the Degree of DOCTOR OF PHILOSOPHY in Biomedical Engineering.

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To my parents Roger J. Guillory Sr., &

Lillie D. Guillory,

And my sisters

Deandrea C. Wright, & Deangela P. Guillory

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Preface

This dissertation is the presented works of 5 interdependent chapters, three of which are published in journals and two in preparation for publication. The contributions of each body of work will be explained rigorously in this section. In all works, Dr. Jeremy Goldman and Dr. Jaroslaw Drelich provided guidance, editing, and final approval of the manuscript. Dr. Jeremy Goldman conducted most surgical in vivo implantations.

The first chapter investigates first-generation zinc materials and relates host inflammatory reaction to corrosion characteristics. It is under the full citation:

1. Guillory, Roger J., Patrick K. Bowen, Sean P. Hopkins, Emily R. Shearier, Elisha J. Earley, Amani A. Gillette, Eli Aghion, Martin Bocks, Jaroslaw W. Drelich, and Jeremy Goldman. "Corrosion characteristics dictate the long-term inflammatory profile of degradable zinc arterial implants." *ACS Biomaterials Science & Engineering* 2, no. 12 (2016): 2355-2364.

The experimental design, execution, writing, and data analysis was mostly performed by Roger J. Guillory II. Dr. Patrick Bowen contributed to SEM and backscattered electron imaging of the implants, and writing/editing sections of the manuscript. Dr. Sean P. Hopkins contributed to data analysis of the macrophage staining and writing. The other co-authors contributed to histological staining procedures and final review of the manuscript.

The second chapter discusses the development of a novel preclinical model to screen degradable metal candidates for cardiovascular stents. The full citation is provided below:

2. Guillory, Roger J., Alexander A. Oliver, Emma K. Davis, Elisha J. Earley, Jaroslaw W. Drelich, and Jeremy Goldman. "Preclinical In Vivo Evaluation and Screening of Zinc-Based Degradable Metals for Endovascular Stents." *JOM* 71, no. 4 (2019): 1436-1446.

The design, execution and data analysis were primarily performed by Roger J. Guillory. Roger J. Guillory wrote the entire manuscript, and all authors contributed to revisions prior to final publication. Emma k. Davis created one of the 7 figures in the manuscript, and Alexander Oliver and Elisha Earley helped with histological sectioning and staining.

The third chapter investigates the relationship between the surface of degradable zinc materials and the subsequent vascular response, using the model described in the second chapter. The full citation is provided below:

3. Guillory II, Roger John, Malgorzata Sikora-Jasinska, Jaroslaw W. Drelich, and Jeremy Goldman. "In Vitro Corrosion and In Vivo Response to Zinc Implants with Electropolished and Anodized Surfaces." *ACS applied materials & interfaces*(2019).

Roger J. Guillory designed the experiments, performed all histology and sectioning, and processed all in-vivo data. Dr. Malgortza Sikora-Jasinska performed all in vitro corrosion experiments, with both authors developing the surface treatments used throughout the study. XPS was performed by Dr. Timothy Leftwich, with sample preparation performed by Roger J. Guillory and Dr. Malgorzata Sikora-Jasinska. Dr. Malgorzata Sikora-Jasinska participated in writing sections of the manuscript and prepared 2 figures fully and one figure jointly. Roger J. Guillory wrote most of the manuscript and prepared 2 figures fully, and one figure jointly.

The fourth chapter describes the newest generation of zinc-based materials developed by our research group. This chapter is in preparation to be submitted to an academic journal. Roger J. Guillory designed the experiments, performed data analysis and wrote the entire manuscript. Dr. Ehsan Mostaed developed the novel material and produced wires from castings. Roger J. Guillory constructed the figures, with the exception of the TEM figure produced by Dr. Ali Mostaed, which was made by Dr. Ehsan Mostaed. The other co-authors participated in histological staining and data collection.

The last chapter investigates an exciting phenomenon seen at the interface of zinc-based materials in the vascular environment. This work is in preparation to be submitted to an academic journal. Roger J. Guillory designed the experiments and performed histological processing, enzymatic assays, data analysis, constructed figures, and wrote the entire manuscript. Alexander Oliver participated in histological data collection and analysis. Timothy Kolesar and Lea Morath performed enzyme activity assays and histological

staining. Dr. Malgorzata-Sikora Jasinska performed ICP analysis from in vitro corrosion studies.

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There are many people, while not contributing to my dissertation directly, that have had substantial influence on me throughout my years here at Michigan Tech. Dr. Tarun Dam,

Dr. Melissa Baird, Susan Libeau, Darnisha Slade Kellie Rafaelli, Dr. Pushpalatha Murthy, Karmen Markham,

Abstract

Biodegradable stents based on zinc have been under development since their introduction in 2013. While metallic zinc is highly ductile, it unfortunately lacks the mechanical strength required for arterial stents. This has led to the development of an abundance of novel zinc-based materials, with the aim of improving the mechanical strength without sacrificing too much ductility. Although these materials are intended to function and slowly degrade within an artery, most zinc-based materials have been developed without deep consideration for their biological effects.

The present work explores the biological effects elicited by zinc-based materials implanted within the arterial system. The biological effects of degradable arterial implants were characterized in terms of quantifiable metrics, including neointimal area, implant to lumen thickness, and base neointimal length. These metrics were used to clarify relationships between material characteristics, including surface oxide film stability, elemental composition, and microstructure, with biological responses. The metrics were also used to compare materials in terms of their biocompatibility. In addition to evaluating biocompatibility, beneficial elements identified by these approaches can be further investigated for their therapeutic value, since all the elements in the implant will be released due to implant degradation. The combined work makes it possible to screen materials in terms of their biocompatibility and provides fundamental insights that impact the metallurgical design of materials.

Introduction

1.2 Heart disease and the state of the art

Heart disease poses an extreme health and monetary risk for Americans. According to the most recent update by the American Heart Association (2017), 92.1 million US adults have at least one type of cardiovascular disease. The average annual cost from heart disease to Americans in 2013 was 199.6 billion USD. Additionally, myocardial infarction and coronary heart disease were 2 of the 10 most expensive hospital diagnoses[1]. The most predominant underlying pathology of myocardial infarction is atherosclerosis. Atherosclerosis is a complex, progressive disease characterized by events such as lipid deposition, intra-arterial plaque formation and buildup, inflammatory mediation and subsequent thrombogenic events emanating from unstable ruptured plaques, all of which contribute to luminal obstruction[2-4].

Presently, the gold standard for treatment of atherosclerosis-induced arterial stenosis is balloon catheter stenting. These metal scaffolds increase the cross-sectional area of the arterial lumen, enhancing downstream blood flow to vital organs (e.g. heart or brain). While the early term effects of stenting diseased arteries are generally positive, the permanent residence of the metal scaffold within the vasculature elicits chronic negative effects. These deleterious effects can include smooth muscle cell intimal hyperplasia due to persistent inflammation[4], stent thrombosis[5], and de novo neoatherosclerosis[2, 6-8].

Over the past 15 years, advances within the interventional cardiology sector have aimed to mitigate early negative outcomes with paclitaxel and sirolimus polymer loaded drug eluting stents[9] (DES), which successfully reduce restenosis rates compared to bare metal stents[10]. Although first generation DESs were effective initially, delayed healing/ endothelialization and an increased risk of very late stent thrombosis have plagued first generation DESs[11]. Second generation everolimus DESs are superior to first generation DESs[10], yet still succumb to

neoatherosclerosis after a long term residence, which appears to be an end term failure mode of any implanted stent regardless of drug elution[7].

1.3 Degradable stents

1.3.1 Polymeric scaffolds

Inadequate mitigation of negative long-term permanent stent outcomes has generated an intense interest for exploring alternative solutions. The most promising alternative approach is to develop a stent that is completely and safely absorbed over time by local vascular tissue. This concept was pioneered by Tami et al, who deployed 25 poly-L-lactic acid stents in 19 lesions across 15 patients with a 6 month follow up time. The study concluded that no significant progressive intimal hyperplasia or lumen loss over 3 and 6 months were provoked by the degradable PLLA based stents[12]. A ten year follow up of the Igaki- Tami stents proved the efficacy of a degradable stent, with mean complete degradation of around 3 years[13]. The 2014 Absorb trials of fully degradable PLLA- everolimus eluting stents showcased the efficacy of drug eluting degradable stents[14]. Although the success of degradable polymeric materials has been clearly shown, this class of materials cannot feasibly achieve the mechanical benchmarks required for arterial stents. This inherent limitation restricts polymeric materials to treating simple lesions of short length. Additionally, the lower radial strength of polymers increases the chances of acute stent recoil and strut malapposition and necessitates an enlarged polymer strut thickness. The excessive strut size and surface area increases arterial injury and thrombogenesis. These serious limitations argue for a degradable metallic alternative.

1.3.2 Degradable metals

1.3.2.1 *Iron*

Peuster and others were some of the first pioneers to investigate iron as a potential degradable metallic stent material [15]. They investigated the efficacy of 16 iron stents in

the descending aorta of New Zealand white rabbits. While these stents performed well with no apparent toxicity and inflammatory response, they did not investigate the ability of the stent to fully corrode. A long term (12 month) follow up study by the authors determined that there was a need to increase the degradation rate of iron materials, as most of the stent remained after the 1 year follow up period [16]. Many investigations after these landmark studies focused on modifications of the bulk iron material to increase the degradation rate [17-25]. More recent absorbable metal candidates have overtaken the use of iron as a base material.

1.3.2.2 *Magnesium*

The most obvious degradable metal candidate, and perhaps the most successful have been materials with a base composition of magnesium. An early study by Heublein and colleagues demonstrated the feasibility of using a non commercial magnesium based alloy to serve as a degradable scaffold, in swine[26]. This initial success for magnesium generated excitement within the degradable metals field, prompting implantations of fully degradable magnesium stents in newborns to treat congenital heart defects, with some degree of success[27, 28]. Unfortunately, the corrosion of magnesium suffers from excessive degradation rates, which is the opposite problem of iron-based materials. With corrosion rates between 19-44 mg/cm²/day, it was estimated that a 5g magnesium implant would lose functionality in 10-20 days, and be completely dissolved within 20-45 days[29]. Magnesium is also known to generate hydrogen gas through the following reaction;

Since magnesium degradation generates around 1L of H₂ per gram of pure material, processing and alloying approaches have received an intense focus from engineers intending to slow the degradation rate of magnesium. A lower degradation rate would be beneficial from an arterial healing standpoint and serve to reduce the rate of potentially

damaging hydrogen gas evolution [30-32]. Although these challenges have precluded the translation of degradable magnesium into safe and effective arterial stents, magnesium still remains the most successful degradable metal material in terms of commercialization progression. A number of clinical trials have demonstrated the efficacy of metal-based degradable stents, including the DREAMS 2G (MAGMARIS) BIOSOLVE II and BIOSOLVE III [33, 34]. Although there has been some success with Mg based scaffolds, problems include mechanical properties that do not yet approach that of conventional metal stents, yielding large strut sizes[35]. Another critical limitation of Mg based stents is the need for a drug eluting coating, which is a known contributor to late term stent thrombosis and delayed endothelialization[36]. Finally, despite extensive metallurgical modifications reported in the literature, Mg stents still degrade too rapidly (<12 months) to support the proper healing of diseased vasculature[35].

1.3.2.3 *Zinc*

Bowen and colleagues introduced zinc in 2013 as a promising degradable metal candidate for coronary stenting[37]. With a penetration rate near the proposed benchmark of 20 $\mu\text{m}/\text{year}$, along with compact corrosion products, zinc materials quickly became a favored candidate for degradable metallic stents[38]. As with the other two degradable metal candidates, the potential upsides come with substantial limitations.

Pure zinc does not possess adequate mechanical properties for stenting, with a tensile strength around 120 MPa, although it has a rather pronounced elongation to failure of 60%-80%[38]. Stenting materials should possess mechanical properties that are approximated in the table below.

Table 1. Mechanical characteristics of Zn compared to stent material benchmark values

Metal	Mechanical Properties			Grain Size [μm]
	Yield Strength [MPa]	Tensile Strength [MPa]	Elongation to Failure [%]	
Benchmark Value	>200	>300	>15	<30
Pure Zn (reference)	<80	<120	60 - 80	>50 - 100

With the introduction of zinc in 2013, questions surfaced regarding the biocompatibility of the metal. Bowen et. al. demonstrated the exceptional biocompatibility of pure zinc within the abdominal aorta of rats[39], using a wire implantation model developed by Pierson et. al. in 2012[40]. Zinc provoked relatively small neointima formations, with an notable gradient of smooth muscle cells receding towards the implant[39]. With the introduction of zinc materials and their optimal biocompatibility, a new wave of material development from metallurgists began to improve the mechanical properties through elemental additions and advanced processing [41-55].

1.4 Scope of dissertation

The excitement generated in the field of degradable Zn based stents led to a bottleneck of novel materials that required biocompatibility evaluations. The in vivo evaluation of degradable materials required a shift in the framework of modern biomaterials development and testing. Our group at Michigan Tech first developed an in-vivo “wire model” to simplify the characterization of the host response to degradable materials

intended for use as cardiovascular stents[40]. This model provides a unique platform to study degradable materials in a fashion that allows detailed examination of the tissue - material interaction. In order to develop successful zinc-based materials, a thorough understanding of the material-arterial cell interactions and the resulting biologic response is essential. We outline these new approaches and investigations in the chapters of this dissertation.

1.4.1 Chapter 1

This chapter investigates multiple zinc materials in an arterial wall implant model. The wall implant model was chosen initially as a simplified environment relative to the luminal space due to the avoidance of blood interaction. This work begins to clarify the relationship between material corrosion character and the ensuing host response. Classically, the foreign body response is a progressive reaction to an implanted biomaterial that eventually leads to fibrotic encapsulation. This classic understanding of the biological response to implanted materials, although widely accepted, must be adapted to incorporate the dynamic and complex biointeractive properties of degradable materials. Understanding the contribution of corrosion products and behavior to the vascular response and overall biocompatibility is of critical importance when designing biodegradable materials.

1.4.2 Chapter 2

Alloying of degradable zinc materials has become a standard approach to improve the mechanical properties. The primary evaluation of materials is a straightforward mechanical testing to compare novel materials to stent material benchmark values. When biocompatibility is evaluated, the primary start point is an insult of corrosion media supernatant extract to 2-D cultured cells. This approach, while useful for understanding basic toxicity, is not appropriate to clarify the in vivo performance of degradable metals. To clarify the biocompatibility of a degradable material, a realistic application environment is an essential start point. Here we develop a quantitative, engineering based

approach to evaluate the performance of degradable materials in an in-vivo system. This approach, while possessing both strengths and weaknesses, can nevertheless help identify the best performing materials using a low cost, physiologically relevant animal model.

1.4.3 Chapter 3

A critical component of material design, particularly in metals that will eventually degrade, are the surface features. Long-term material corrosion characteristics of zinc materials can be directed by the material's initial surface conditions. Since the potential exists for the control of surface properties to make possible the "tunability" of corrosion dynamics, the relationship of surface characteristics to the vascular response must be clarified. This chapter describes an investigation to relate varied surface features of degradable zinc-based implants to the vascular response.

1.4.4 Chapter 4

Our group at Michigan tech has steadily developed a suite of degradable metal candidates for cardiovascular stents. Previously, it has been challenging to relate differences in neointimal character to specific modifications of materials. This chapter evaluates a new generation of degradable zinc alloys and provides insight into the relationship between elemental compositions and neointimal formation. Understanding the interplay between metallic elemental additions and vascular responses is critical to designing top performing alloys. In parallel, we have also discovered that changes to the microstructure affect neointimal responses. These two components of alloy design not only alter the mechanical properties of materials, but also critically impact biological outcomes.

1.4.5 Chapter 5

To date, we have examined zinc materials from a performance standpoint in order to select the best performing materials for continuation in the stent development pipeline. While this approach is useful from an engineering and commercialization standpoint, there is tremendous value in understanding the basic interactions between the material

and arterial cells. Recently, Bowen and colleagues described an exciting phenomenon of neointimal smooth muscle cells organizing peripherally from the degradable zinc implant and avoiding the interface. In this contribution, the neointima of pure zinc materials is probed for smooth muscle cell content and compared to a non-degrading control. Molecules involved with the regulation and initiation of apoptosis are monitored. A material that can exert an anti-proliferative/pro-apoptotic effect on the migratory smooth muscle cells is ideal within the realm of vascular materials. Current generation stents require a drug eluting polymer coating to prevent excessive neointimal growth and limit harmful outcomes. Here, we find that the byproducts of zinc implant degradation may impart a suppressive effect against vascular smooth muscle cells and anti-inflammatory pro-healing effects on endothelial cells. This paradigm shift that emerges with degradable zinc materials within the vasculature may shift the way scientists think about degradation as a whole; not just as a peripheral feature of the material, but as central means of imparting continuous therapy to the treated vessel for the entire lifetime of the implant.

1.5 Summary

This dissertation provides new insights into the biocompatibility of degradable zinc implants for use in the cardiovascular system. Chapter 1 provides a rigorous and detailed histopathological analysis of zinc and zinc alloys in an in vivo implant model. The methodology used allows for a multifaceted approach to understanding the dynamic foreign body response to degrading materials. From this work, we are able to determine that the inflammatory response not only depends on the elemental additions, but manifests primarily from the inherent corrosion characteristics dictated by the elemental additions. These results demonstrate that tissue reactions towards degrading zinc materials can be regulated by controlling corrosion behavior.

With this newfound knowledge, a natural hypothesis arises that different degradable zinc materials (ie. Zn-Li vs. Zn-Mg) with different inherent corrosion properties will evoke different responses in the vascular system. To test this hypothesis, an analysis framework with appropriate design and outcome metrics needs to be established. Chapter 2

accomplishes this by developing a reliable method for the quantification of biocompatibility metrics within the rodent aorta. This standardized methodology makes it possible to compare materials in terms of biocompatibility and relate the differences in biocompatibility of materials to material characteristics. By using histomorphometry parameters that are similar to those used when evaluating stents in the clinic, the results from the characterization of neointimal responses to wire implants can be compared to stents that are in clinical use.

Since it is apparent that corrosion characteristics affect neointimal growth and development, variations in the initiation of corrosion would be expected to dramatically alter the vascular response. In chapter 3 we vary the surface oxide film character of pure Zn substrates to investigate the effect of surface oxide film properties on early term neointimal responses. Interestingly, standard surface treatments used for permanent metallic materials may not be optimal for zinc implants. This is due to the initiation of harmful corrosion events that exacerbate neointimal progression when using zinc implants with electropolished surfaces.

Chapter 4 probes deeper into the neointimal responses of a wide variety of new generation alloys, using the established framework presented in Chapter 2. Complex alloy systems (elemental additions >3) provide unique changes at a microstructural level, which are found to directly impact the neointimal response/ performance of the material. Understanding the relationship between the microstructure and neointimal response provides predictive power that aids in the design of future materials.

Even though this body of work rigorously demonstrates that bulk corrosion, surface characteristics, and microstructural changes in Zn materials impact biocompatibility, the role of the corrosion byproducts in the biocompatibility of the implant remains a critical open question. Chapter 5 probes the cellular and protein content in the neointimal tissue surrounding Zn based implants. The results of the analyses demonstrate that ionic Zn initiates apoptotic death in vascular smooth muscle cells. Importantly, this chapter shows that metallic zinc may serve not only as structural support for the vessel, but its release in

an ionic form profoundly affects neointimal cells through its interaction with protein-mediated cell signaling pathways.

Together, this body of work evaluates the impact of zinc-based materials as potential cardiovascular stents on biocompatibility by integrating a large spectrum of fields ranging from materials science, physiology, and cell and molecular biology. During the course of this multifaceted investigation, novel approaches are developed and material properties are related to neointimal growth and device performance.

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1 Understanding zinc material corrosion characteristics and the foreign body response progression in the arterial environment ^{1 2}

1.1 Introduction

The concept of constructing fully bioresorbable stents (BRSs) to open occluded arteries using non-toxic degradable materials has inspired extensive research over the past two decades, with an emphasis placed on polymer-based materials ¹. While multiple polymer-based fully degradable BRS have proceeded through early clinical trials in the US and are beginning to obtain FDA approval², the development of stents manufactured from metallic materials has largely been restricted to pre-clinical studies ^{1b}. This has been the case despite the order-of-magnitude mechanical strength advantage that metals derive over polymers and their similarity to conventional stents in terms of deployment and mechanical characteristics.

Magnesium has been extensively investigated for endovascular stenting and is the only biometallic stent studied in the clinical setting. The benefits of magnesium-based stents include low thrombogenicity, excellent biocompatibility to neighboring cells and tissue, and good target lesion failure rates at 6 months, especially when coated with an antiproliferative drug ³. However, early generation magnesium stents were found to have high degradation rates and even the 2nd generation stents have late lumen loss rates much higher than standard DESs and some of the polymer-based BRSs ⁴. Though studied less in recent years, iron-based materials have the favorable profile of producing low to moderate inflammatory responses and possess excellent mechanical properties ⁵, but produce voluminous corrosion products that build up in the artery and repel neighboring cells and biological matrices ⁶. The serious limitations of iron and magnesium and the intense focus on surmounting them with incremental material modifications have delayed

¹ This chapter contains material previously published in ACS Biomaterials Science and Engineering

² References in Section 1 follow the ACS author guidelines.

developmental efforts for bioabsorbable metallic stents relative to polymers. Recently, zinc and its alloys were introduced as stent materials that do not experience the limitations seen with iron and magnesium⁷. Zinc could potentially break through the considerable developmental hurdles facing absorbable biometals (for a recent review of zinc and its progress towards becoming a candidate material for endovascular stents, refer to ^{1b}).

In contrast to an inflammatory response generated around a biostable material, which is directed primarily against the base material—metals and their stable oxide layer in the case of bare metal stents, or polymers in drug-eluting variants—the inflammatory response to a corrodible metal may also be impacted by the corrosion behavior and the products thereby generated. To date, most studies have focused on the *in vitro* degradation and cytotoxicity of zinc and zinc alloys⁸. With the exception of limited-in-scope *in vivo* work performed by our group ^{1b, 7, 9}, a detailed investigation into the cellular and tissue interactions with metallic based corrosion products for a degradable metal remains largely unexplored ^{1b}. Due to the complex biochemical and cellular constituents present in the physiological environment that impact the generation and progression of corrosion products, as well as their integration and ultimate break down and clearance, a long-term *in vivo* evaluation is essential to clarify the biological response.

The present work was undertaken to characterize the chronic inflammatory response to zinc by drawing detailed comparisons between different zinc-based materials displaying a range of corrosion behaviors. We describe the cell/tissue remodeling events that take place on the active tissue-material interface of degrading zinc implants in order to understand the inflammatory progression. Through a direct comparison it may be possible to gauge the inflammation response severity and clarify how corrosion behavior may regulate inflammatory pathways. This type of analysis may eventually allow for an improved design of zinc-based stent materials with the goal of minimizing the severity of chronic inflammatory responses.

1.2 Experimental

1.2.1 Materials and reagents

0.25 mm diameter wire of 99.99+ wt.% (4N) zinc was purchased from Goodfellow Corporation. Zn-Al alloy ingots were purchased from Metropolitan Alloy Co. (Detroit, MI) at concentrations of 0, 1, 3, and 5.5 wt. % Al. The Zn stock used for fabrication was special high grade (SHG) ~99.7% zinc per ASTM B6-13. Ingots were gravity cast, in air, into flat steel molds. The pancake ingots received measured approximately $100 \times 70 \times 10$ mm. Alloying with aluminum substantially improved the mechanical properties of the zinc material (the detailed microstructure and mechanical properties characterization are included in a separate contribution¹⁰). Table 1-1 provides the list of impurities and their concentrations in both 4N and SHG Zn.

Table 1-1 – Impurity levels in 4N and SHG Zn

Impurity	Concentration (ppm)	
	4N ^a	SHG ^b
Ag	5	< 2
Al	n/s	< 2
As	n/s	< 2
Cd	7	< 2
Cu	5	7
Fe	25	371
Mg	n/s	< 2
Mn	< 1	3
Pb	15	16
Si	< 1	n/a

^a Per specification; non-specified impurities are labeled “n/s”

^b Measured via inductively coupled plasma optical emission spectroscopy; non-analyzed impurities are labeled “n/a”

Electro-discharge machining was used to fabricate SHG and Zn-Al strips measuring approximately 12 mm long $\times$ 300 μ m wide $\times$ 300 μ m thick from the rolled foil. The

strips were electropolished using a stirred bath of 120 mL deionized water, 885 mL ethyl alcohol (200-proof – Pharmco Aaper, Brookfield, CT), 100 mL n-butyl alcohol (99.9% - Fisher Scientific, Fair Lawn, NJ), 250 g anhydrous zinc chloride (reagent grade – Avantor, Center Valley, PA), and 109 g aluminum trichloride hexahydrate (97+% - Sigma Aldrich, St. Louis, MO) at a constant potential of 10 V for 90 seconds ¹¹. Due to the electropolishing, the thickness of strips was reduced to 220-270 μ m and their rectangular cross-sectional dimension aligned closely with that of a wire and are hereafter referred to as wires.

Staining reagents included eosin Y disodium salt, acetic acid (99.7%), toluidine blue (powdered), phosphotungstic acid solution 10% w/v, phosphomolybdic acid hydrate solution, biebrich scarlet acid-fuchsin solution, analine blue solution, absolute ethanol, xylene substitute, Gill's No. 3 hematoxylin solution, hematoxylin (powdered), anhydrous iron (III) chloride (powder 99.99%), Lugol's iodine Solution phosphate buffered saline (PBS), phosphate buffered saline with bovine serum albumin (PBS-BSA), and Eukitt quick-hardening mounting medium; all were obtained from Sigma Aldrich. Hydrochloric acid (HCl) (37%; 12.2 M) was obtained from EMPROVE (Darmstadt, Germany).

Reagent grade acetone was obtained from Pharmco-Aaper.

Primary antibodies included anti-CDd68 (ab125212), anti-Cd11b (ab8879) all purchased from Abcam (Cambridge, Massachusetts). Secondary antibodies include goat anti-rabbit IgG alexa fluor 488 (ab150077), donkey anti-rabbit IgG alexa fluor 555 (ab150074) purchased from Abcam, and anti-CD163 (sc-58965) purchased from Santa Cruz Biotechnology (Dallas, Texas). Streptavidin alexa fluor 633 conjugate (S-21375) was purchased from Life Technologies (Carlsbad, California). Biotinylated goat anti rabbit IgG (ab64256), biotinylated goat anti mouse IgG, and endogenous avidin/biotin blocking kit (ab64212) were all purchased from Abcam. Goat serum (G9023), 4', 6-diamidino-2-phenylindole dihydrochloride (DAPI), Triton X-100, phosphate buffered saline (pH 7.4 containing 1% BSA w/v) were all purchased from Sigma-Aldrich. PermaFluor mountant was purchased from Thermo-Scientific (Waltham, Massachusetts).

1.2.2 In vivo implantation and cryo-sectioning

The zinc implant wires including 4N, SHG, Zn-1Al, Zn-3Al, and Zn-5Al were cleaned in acetone and then sterilized in 70% (v/v) ethyl alcohol for 20 to 30 min prior to implantation. A surgical protocol developed recently was employed to implant two wires of the same material composition into the abdominal aorta wall of an adult Sprague-Dawley rat⁶. Briefly, the wires were used to penetrate the adventitial layer of the abdominal aorta, after which the wire was advanced within the arterial wall for the full length of the specimen. We have found that the wall environment is more conducive to biocorrosion activity whereas blood contact from a luminal implantation tends to passivate the metal surface and reduce corrosion. Thus, the wall implantation was employed to increase the bioactivity of the metal surface and elicit the very inflammatory responses that we were interested in investigating. The actual inflammatory response to a stent made from the same material may therefore be less severe than what has been found in the present study. The diameter of the wires is approximately twice that of a stent strut of the same material, thereby introducing a greater challenge to the local arterial environment in terms of material degradation and corrosion product clearance. The SHG, Zn-1Al, and Zn-3Al alloy wires were extracted at four time points, including 1.5, 3, 4.5, and 6 Months. The Zn-5Al alloy wires were extracted at 1.5, 3, 3.5, and 6 months. The 4N wires were extracted at 13 time points, including 1–10, 12, 15, and 20 months. One of the two wires implanted into each rat was used for metallographic and corrosion rate analysis, while the second wire was used for inflammatory characterization work (n = 28 rats). All animals survived the wire implantation surgery and made a full recovery. All animal experiments were approved by the animal care and use committee (IACUC) of Michigan Technological University.

At the indicated time points, wires were excised for histological evaluation and immunofluorescent labeling by collecting the entire segment of host artery that contained the implant. To facilitate cross-sectioning, the lumen of the collected artery was filled with PolyFreeze mounting medium (Sigma Aldrich) by syringe perfusion. Explanted arteries were placed into plastic molds, surrounded with PolyFreeze mounting medium, and snap-

frozen in liquid nitrogen. Samples were then stored at -80°C until cryo-sectioning. Frozen samples were cross-sectioned at a 10 µm thickness with a Thermo-Scientific HM 550 P cryostat. Tissue sections were placed on warm VWR Histobond (Radnor, PA) slides and fixed in -20°C absolute ethanol for 60 s or in 4% formaldehyde (Sigma Aldrich) and washed three times in PBS (Sigma Aldrich) for 5 min prior to staining.

1.2.3 SEM preparation and backscattered imaging

A JEOL JSM-6400 (Peabody, MA) scanning electron microscope equipped with a dSpec automation system (Geller MicroAnalytical Laboratory; Topsfield, MA) was used for examining the explant cross sections. Imaging of the carbon-coated specimens was conducted at 10 kV accelerating voltage at a reduced working distance using a backscattered electron detector. The acquired backscattered electron images were analyzed with imageJ (National Institute of Mental Health; Bethesda, Maryland) to yield cross sectional area measurements. The bright zinc portion of the image was selected by thresholding, in which only the area containing the brightest pixels was measured. From these cross sectional area measurements, a metal cross sectional area reduction was calculated for each image. The resulting measurements were averaged to yield an estimated area reduction.

1.2.4 Hematoxylin and eosin

Slides with fixed tissue sections were rinsed in deionized water for 5 min to clear any residual PBS. Gills-3 hematoxylin solution was pipetted onto each slide for 2-5 min, or until over-stained (determined by visual examination). Slides were then dipped in a 900 mL bath of distilled water that had been acidified to pH = 1.8–2.0 by adding HCl. After 5 dips in the acidic bath, the samples were rinsed in distilled water for 1-5 min. Slides were then washed twice with 95% ethanol (v/v). The slides were counterstained with Eosin Y working solution (0.25%), which was micro-pipetted onto each slide for 45 s. immediately after counterstaining, the slides were rinsed and dehydrated in two changes of absolute ethanol for 5 min apiece. The slides were then cleared twice with xylene

substitute for 5 min, and mounted under a cover slip using Eukitt mounting medium. Samples were imaged using an Olympus BX51, DP70 bright-field microscope (Upper Saucon Township, PA).

1.2.5 Toluidine blue

A Toluidine Blue stock solution was prepared by dissolving 1 g of toluidine blue in 70% ethanol. A working solution was prepared by using 5mL of the stock solution and 45mL of a 1% sodium chloride solution with the pH adjusted to 2.4 using HCl or NaOH. The slides were then hydrated in three changes of DiH₂O for 5 min each and stained in the working solution for 5 min. The slides were then dipped 3 times in 95% ethanol and 5 times in 100% ethanol to quickly dehydrate. Slides were then dipped 20 times in each of two clearing solutions—a 1:1 xylene substitute/acetone and a pure xylene substitute mixture—mounted in resinous mounting medium, and imaged with an Olympus BX51, DP70 bright-field microscope.

1.2.6 Immuno-fluorescence

Biological tissue surrounding the metal implant became impregnated with an auto-fluorescent corrosion product. To remove the background fluorescence due to the corrosion product and improve the overall quality of immuno-fluorescent labeling, slides were immersed in a 1% (v/v) acetic acid solution for 1 h to dissolve the corrosion product. Slides were then rinsed in 4 changes of PBS for 5 min each and incubated in a 0.2% Triton X-100 solution in PBS for 30 min, followed by immersion in PBS-BSA and a 0.2% Triton X-100 solution (blocking buffer) for 30 min. If biotinylated secondaries were used, avidin/biotin blocking was performed for 15 min each with the endogenous avidin/biotin blocking kit purchased from Abcam, as well as the appropriate IgG. Primaries were then incubated overnight on the slides at 4°C in a 1:100 dilution, followed by 3 washes in blocking buffer for 5 min each. For CD163 and CD11b antibodies, a concentration of 1:500 was used. Secondaries were incubated on the slides at room temperature in 1:200 concentrations for 1 h. Three washes in blocking buffer were

performed for 5 min each and the slides were incubated with a DAPI working solution for 1 min (1 μ l/mL DAPI solution). Slides were then rinsed in 2 changes of PBS and mounted with Perma-Fluor aqueous mounting medium and imaged with appropriate fluorescent filters.

1.2.7 Immunofluorescent Quantification

For quantification of immunofluorescent staining, a minimum of 3 tissue sections were taken of each sample separated by at least 100 μ m, and stained with the aforementioned procedures. Sections were then imaged using the appropriate filters at 100X normal magnification, and uploaded to MetaMorph imaging software. The entire inflammatory exudate was included in the area selected for automatic pixel counting. Data was recorded and graphed using Excel 2013.

1.3 Results

1.3.1 Backscattered imaging and cross sectional area reduction calculations

Figure 1-1. Backscattered imaging and cross-sectional area reduction calculation for the five evaluated zinc metal compositions, 4N wire and the four strip materials. The backscattered images shown are for the 5 compositions at 1.5 months post-implantation, while the cross sectional area reductions shown in the graph include samples at 1.5, 3, 4.5, and 6 months.

The SHG, Zn-1Al, Zn-3Al and Zn-5Al metal wires had the same surface finish after electropolishing, with oxide film thickness between 100 and 200 nm. Only the 4N wire had a native oxide film, produced during wire manufacturing, but this film had a similar thickness as presented earlier¹². Quality of the surface finish influences the early stage of

biodegradation in the first few weeks after implantation as discussed in the cited reference, but has no effect on corrosion of the Zn implant in later stages, as discussed in this contribution.

Fig. 1-1 shows examples of metallographic cross sections of 1.5-month specimens. Metallic portion of the explant is represented by the brightest area whereas oxidized metal is in a grey color. These metallographic cross sections revealed sharply varying corrosion behavior between the five zinc compositions that were evaluated. The 4N wire experienced a high resistance to corrosion at this time. The SHG experienced a similar mode of corrosion, with a significantly higher cross sectional area reduction of metallic portion relative to the 4N material. The Al alloys experienced a similar cross sectional area reduction as the SHG, but the surface corrosion mode shifted to intergranular attack.

1.3.2 Hematoxylin and Eosin

Fig. 1-2. Representative H&E stains of cross-sections containing the explanted artery with the 4N wire implant. Panels A-H represent months 1, 2, 4, 5, 6, 8, 9, 20, respectively. The green asterisk in each image denotes the wire or corrosion layer location. The scale bar at lower right is 100 μ m.

The inflammatory response progression against the 4N wire implantations from 1 to 20 months is shown in Fig. 1-2. Panel A shows a small amount of cellular alignment around the metallic implant with mononuclear infiltrates into the surrounding tissue and negligible corrosion of the metallic implant. Panels B and C (2 and 4 months) show progression towards a dense, non-porous corrosion product occupying the space at the original interface between biological tissue and metallic wire, as well as an increase in cellular density at the material interface. Panels D and E (5 and 6 months) show areas near the interface of the implant that have become devoid of nuclei and contain irregular nuclear features. Irregular nuclear features decrease dramatically in the peripheral tissue, with no abnormalities in the media or intima of the artery for each time point. Panels F, G and H (8, 9, and 20 months respectively) depict an increase in fibrotic deposition around the implant, as well as localized chronic inflammation inside the fibrotic capsule and around the interface. Increased image size and identification of important cells/tissue remodeling events are shown in Fig.s1-S3 and 1-S4.

Fig. 1-3. Representative H&E stains of cross-sections containing the explanted SHG, Zn-1Al, Zn-3Al, and Zn-5Al wire implants within the arterial wall over 1.5 – 6 months, and Zn-5Al at 1.5, 3, 3.5, 6 months post implantation. The green asterisk denotes the wire or corrosion layer position. The scale bar at lower-right is 100 μ m.

The progression of the inflammatory response against SHG, Zn-1Al, and Zn-3Al, implants at 1.5, 3, 4.5, and 6 months, as well as at 1.5, 3, and 3.5 months for Zn-8Al is shown in Fig. 1-3. A large corrosion layer is observed for the SHG specimens at 1.5 and 3 months, replete with macrophages and an irregular material - tissue interface. An increase in flocculent corrosion product infiltrated with viable cells was identified on H&E stains for SHG at 4.5 and 6 months. Tissue deposition is within the original collagenous capsule at 6 months. An increased amount of corrosion relative to 4N and a more intense cellular infiltration relative to SHG is observed within the corrosion layer for Zn-1Al at 1.5 months. At 3 months, an increase in capsule density is seen for Zn-1Al

in Fig. 1-3, as well as infiltrating cells with fragmented nuclei. For Zn-1Al at 4.5 months, an increase in fibroblasts and cellular organization is evident, with active chronic inflammatory cells present inside the corrosion layer. A highly organized collagenous capsule is apparent at 6 months for Zn-1Al, with neovascularization and localized chronic inflammation. For the Zn-3Al series of implants, a high degree of corrosion and cellular infiltration with an ill-defined tissue material interface at 1.5 months is shown in Fig. 1-3. At 3 months, a highly organized and cellularized collagenous capsule formed around Zn-3Al, with fibroblasts present at the interface and inflammatory cells inside the interface and corrosion layer. Some nuclear fragmentation is also visible in the cells present within the corrosion layer. At 4.5 months for Zn-3Al, an increase in nuclear disparity was noticed inside the corrosion layer. Collagen bundles of fibrotic character surround the implant, with an increasing amount of acellularity relative to the 3 month specimen. At 6 months for Zn-3Al, a dense collagenous capsule with sparse amounts of cells surrounds the implant. Macrophages still impregnate the corrosion layer and an active tissue-material interface is still evident. The Zn-5Al 1.5 month specimen displayed the most intense initial inflammatory reaction of all the specimens, with a high cell density at and around the material tissue interface. At 3 months, concentric cellular alignment was formed around the corrosion layer interface, which increased in its regularity at 3.5 months. A dense, fibrous capsule was built around the corrosion layer for the 3.5 month time point. For more detailed descriptions, see Figs 1-S5-1-S8.

1.3.3 DAPI Staining

Fig. 1-4. Representative DAPI images showing corrosion layer and tissue interface for 4N, SHG, Zn-1Al, and Zn-3Al implants at 1–1.5 and 6 months. The green dashed line identifies the interface between corrosion layer and tissue. The yellow dashed line identifies the location of the old interface. The scale bar at lower-right is 100 μ m.

Because it was sometimes challenging to clearly visualize cells present within and near the corrosion layer in the H&E stains, DAPI staining was used to help improve the nuclear characterization of this cell population for the different zinc materials.

Representative images for each zinc composition, excluding Zn-5Al, are shown at approximately 1–1.5 months and 6 months (Fig. 1-4). The 4N material exhibited a slight presence of intact mononuclear cells directly at the rigid interface at 1 month. At 6 months, a dense and uniform population of both intact and dying cells was present within the corrosion layer. Dying cells are evident throughout this layer by the appearance of nuclear fading, shrinkage, and fragmentation. Cells in the surrounding tissue are highly aligned. For SHG zinc at 1.5 months, a dense and uniform population of both intact and dying cells was present within the corrosion layer. Surrounding cells displayed some degree of alignment as well as the general absence of inflammatory cells. At 6 months,

the corrosion layer-tissue interface appeared to move closer to the zinc implant, potentially indicative of a dynamic interface. The space between the new and old interface was uniformly populated with viable cells, indicative of healthy tissue regeneration having taken place. At 1.5 months for Zn-1Al and Zn-3Al, a high density of mononuclear cells is present immediately adjacent to the tissue interface with the corrosion layer, symptomatic of an intense inflammatory response in the surrounding tissue. The coverage of cells within the corrosion layer was not uniform, with open spaces evident at numerous locations, revealing an environment that may be toxic to cells. These cells displayed a greater degree of intense nuclear staining relative to what was seen surrounding the SHG material, consistent with nucleic acid condensation ¹³. At 6 months, the interface had advanced closer to the implant. However, the space between the old and new interface was largely devoid of cell nuclei, symptomatic of poor tissue regeneration. In general, the results indicate moderately healthy tissue regeneration around the SHG implant in contrast to toxicity in combination with poor tissue regeneration around the Zn-1Al and Zn-3Al implants due to apparent differences in the inflammatory response. Greater detail is presented in Fig.s 1-S9 and 1-S10.

1.3.4 Toluidine Blue

Fig. 1-5. Quantification of average number of mast cells via multiple sections per sample.

Mast cell counting via toluidine blue staining was performed for all alloy compositions up to 6 months as shown in Fig. 1-5. A slightly higher mast cell average is seen for the initial months of the SHG specimen, relative to the Al alloys, with a slightly increasing trend over a period of 1 to 6 months. The average number of mast cells counted for the Zn-Al alloys increased more sharply over 6 months compared with the 4N and SHG material, revealing a temporal increase of inflammatory intensity for the Zn-Al alloys. Average mast cell counts for the SHG specimen at 6 months ($\pm 11 \pm 6$) is slightly lower when compared to 4N at the same time ($\pm 16 \pm 5$). The Zn-Al alloys' average mast cell

count after 6 months was $\pm 19 \pm 5$, $\pm 20 \pm 3$, and $\pm 6.526 \pm 7$ for 2Al, 4Al, and 8Al respectively. The Zn-Al alloy counts were all higher than for the SHG and 4N samples.

Fig. 1-6. Toluidine Blue staining of 4N (panels A, B), and SHG (panel C) implants at 6 months. The image shown in panel A is from a region in close proximity to the implant. The image shown in panel B is from the peripheral tissue, approximately 1 mm from the implant surface. Yellow arrows denote mast cells. The scale bar in the lower image is 100 μ m.

Toluidine Blue staining revealed the presence of active (degranulating) mast cells near the 6 month 4N implant (Fig. 6, panel A) and in the surrounding tissue (panel B). Panel C in Fig. 1-6 shows a positive but inactive (quiescent) mast cell staining near the 6-month

SHG implant. All mast cells (degranulating and quiescent) were included in counting for completeness.

1.3.5 CD68,CD11b, and CD163 fluorescence

Fig. 1-7. CD68 (green) and CD11/b (red) labeling of cross-sections containing the explanted artery and the 4N, SHG, Zn-1Al, Zn-3Al and Zn-5Al compositions at 6 months. Cell nuclei were counterstained with DAPI (blue). High magnification CD68 images at the tissue-material interface and in the peripheral tissue are shown in the 2nd and 3rd rows for each specimen. The scale bar for the 100× nominal magnification shown at right (top row) is 500µm. Background auto-fluorescence of the elastin rich media layer is seen in the Zn-5Al specimen. The scale bar for the 600× nominal magnification images shown at right (3rd row) is 100µm. The scale bar for the 200x nominal magnification images shown at the right of the bottom row is 200µm.

Fluorescent antibody labeling was performed to identify the response of CD68⁺ and CD11b⁺ cells to the different zinc compositions (Fig. 1-7). The CD68 glycoprotein is highly expressed in monocyte/macrophage populations (M ϕ)¹⁴ A large population of CD68⁺ cells surrounded both the 4N and Zn-3Al implants. A thick CD68⁺ cell layer was established at the interface of these two materials, although the positive cell layer at the Zn-3Al interface was more continuous and uniform relative to a more irregular 4N interface that did not surround the entire metal specimen. The media layer in the Zn-3Al specimen showed intense positive staining for CD68. A more detailed examination revealed substantial damage to the ECM of the media layer in the Zn-3Al section (data not shown). In contrast, the SHG and Zn-1Al compositions exhibited moderate CD68 labeling at the interface and sharply decreasing numbers of positive cells in peripheral tissues, although a modest increase in positive labeling was apparent in the Zn-1Al periphery relative to SHG.

Fig. 1-8. Quantification of CD68 staining using average pixel counting of multiple tissue sections per sample.

Trends of relative CD68 expression for each sample was evaluated and quantified (Fig. 1-8). Interestingly, the 4N CD68 expression at the 1 month time point, $(2 \pm 1) \times 10^4$, $\times 10^4 \pm \times 10^4$ is substantially lower than the SHG response at 1.5 months, $(10 \pm 3) \times 10^4$, $\times 10^4 \pm \times 10^4$ with increasing vs. decreasing counts respectively.

CD11b and CD163 labeling was performed on all specimens and quantified up to 6 months (Fig.s S1 and S2). Cells that express CD11b are mostly granulocytes, such as neutrophils. Macrophages can also express CD11b, but to a reduced extent relative to

neutrophils. The presence and distribution of CD11b⁺ cells provide fundamental information related to the stage of inflammation at a wound site. In our 6 month samples, CD11b⁺ cells were present near the interface of the 4N and Zn-3Al compositions, while the SHG and Zn-1Al compositions experienced a reduced presence of CD11b positive cells both near and far from the interface. CD163⁺ cells were believed to be alternatively activated macrophages, commonly known as M2 macrophages, and generally decrease the inflammatory response around a wound site. They also encourage tissue repair by releasing urea, polyamines, and ornithine¹⁵. Tissue with a more resolved state of inflammation around an implant will contain CD163⁺ cells, while a chronic inflammatory response will most likely have little to no CD163⁺ cells¹⁶. In our 6 month samples, CD163⁺ cells were seen very far away from the 4N implant, but not near the interface. The SHG and Zn-1Al implants were the only implants to show CD163⁺ cells at the tissue-implant interface. The Zn-3Al implant was largely void of CD163⁺ cells.

At 6 months, the 4N and SHG materials elicited a similar expression level (positive pixel count, Fig. 1-S1) of CD163 labeling, $(4 \pm 2) \times 10^4$ and $(3 \pm 2) \times 10^4$, respectively. $\times 10^4 \pm \times 10^4 \times 10^4 \pm 2.4 \times 10^4$ This was the case, although they demonstrated dissimilar early time point expression values, $(0.5 \pm 0.3) \times 10^4$ and $(4 \pm 2) \times 10^4$, respectively. $\times 10^3 \pm \times 10^3 \times 10^4 \pm \times 10^4$ The Zn-1Al, Zn-3Al, and Zn-5Al alloys elicited relatively low expression levels of CD163 at 6 months, $(1.6 \pm 0.9) \times 10^4$, $(1.2 \pm 0.3) \times 10^4$, and $(2.4 \pm 0.5) \times 10^4$ respectively. $\times 10^4 \pm \times 10^3 \times 10^4 \pm \times 10^3 \times 10^4 \pm \times 10^3$

According to Fig. 1-S2, the Zn-1Al, Zn-3Al, and Zn-5Al alloys elicited an initially high CD11b presence, $(5 \pm 1) \times 10^4$, $(4 \pm 1) \times 10^4$, $(5 \pm 1) \times 10^4$, $\times 10^4 \pm \times 10^4 \times 10^4 \pm \times 10^4 \times 10^4 \pm \times 10^4$ when compared to the 4N and SHG materials, $(3 \pm 1) \times 10^4$ and $(1.8 \pm 0.5) \times 10^4$, respectively. $\times 10^4 \pm \times 10^4 \times 10^4 \pm 4.7 \times 10^3$

1.4 Discussion

An interesting finding in this work is that the inflammatory response evolution differed dramatically between five zinc-dominant materials with minor additions of non-zinc impurities or alloying elements. The compositional impurities that ranged from ~0.01 (4N) to ~0.3% (SHG) produced dramatic differences in inflammation. This was the case despite a similar cross sectional area reduction by 6 months for all examined materials. Surprisingly, the highest purity 4N zinc implant elicited the strongest negative immune response. Minor impurity introductions of 0.3 wt.% (SHG) produced the best performing zinc implant of the five, with a highly biocompatible immune response. The results demonstrate that the immune response against zinc implants was not related to zinc *per se*, but is rather shaped by the material's intrinsic corrosion behavior. Because zinc 4N corrosion rate at early times was substantially lower than that of SHG zinc, we speculate that the initial corrosion rate may be a critical regulator of inflammation.

An analysis of the histological and metallographic data provides insight into the potential mechanism by which the corrosion rate may regulate the inflammatory response. For the 4N composition, a necrotic response at 5, 6, and 8 months (Fig. 1-2, panels D–F respectively) is highly indicative of inflammation-induced matrix remodeling around the implant. The necrosis is believed to have occurred due to the generation of a mature fibrotic capsule between 4 and 6 months, coupled with what appear to be intense, localized corrosion events that damage the fibrotic capsule. We have reported that the stable ZnO layer present around the 4N implant increases into a bulk corrosion product at 4.5 and 6 months ⁹. The temporal trend of necrosis appearing by 5 months and a bulk ZnO product appearing by 4.5 months is associated with our recent report that the 4N zinc corrosion rate accelerates between 4.5–6 months ⁹. Recent electrochemical and thermodynamic characterization of zinc corrosion as a function of pH ¹⁷ has identified ionic zinc as the most stable species of zinc at low pH conditions (pH ~ 1- 4) Zn^{2+17} , while ZnO is the most thermodynamically stable phase of zinc at physiological pH. The presence of necrotic tissue at the 4N interface (Fig. 1-2) at the time points where we

found an increased corrosion rate [8]. Such an environment could have materialized if frustrated phagocytosis by foreign body giant cells (FBGCs) and resident Mφs at the 4N implant lowered the interfacial pH to acidic conditions. The resulting ionic Zn^{2+} release could induce massive cell death near the implant surface, due to a low reported cell tolerance to ionic zinc ^{8a, 8d}, and stimulate fibrotic tissue deposition. This putative mechanism may explain the widespread cell death seen at 5, 6 and 8 months and the ensuing fibrotic capsule growth near the surface of the 4N material, at which time points we have also found a high CD68 and CD11b signal. Similar reasoning could also explain the lack of cell death in the SHG material. In this case, downward temporal trending of CD68+ Mφ (Fig. 1-8) around the SHG implant may reflect a near neutral pH interfacial environment and effective Mφ clearance of porous ZnO and ZnOH corrosion products.

It is well known that foreign particle size and scaffold porosity affect Mφ signaling, phagocytosis, and fusion ¹⁸. We speculate that the initial low corrosion rate of the 4N material may have elicited macrophage fusion (observed, but not shown) by presenting a rigid interface and eventually large corrosion particles or a low porosity scaffold to the surrounding tissue. Mφ fusion / foreign body giant cells can cause tissue damage by releasing powerful enzymes and reactive oxygen species in an effort to degrade the material into smaller, digestible particles ¹⁹. Interestingly, the 4N corroding implant elicited the degranulation of mast cells (Figs 1-5 and 1-6), The findings of inflammatory events at 6 months for the 4N material that are normally associated with acute inflammation (Fig. 1-5) are consistent with the concept of an increased Mφ activity that accelerates corrosion and increases toxicity to cells near the implant surface.

In contrast, the SHG and Zn-Al metals all appear to degrade earlier than the 4N material and produce a relatively thick and porous corrosion layer (Fig. 1-3). This also correlates well with temporal CD68 expression, as presented in Fig. 1-8. The porosity is apparent due to the increased consistency of cellular infiltration within the corrosion layer for SHG and the Zn-Al alloys. The ability of inflammatory cells to penetrate the corrosion layer at early times may explain the reduced thickness of the fibrous capsule in SHG relative to 4N. Interestingly, the 4N material was the only implant to elicit an acellular fibrotic

encapsulation response, despite the similar material compositions and preponderance of zinc amongst the SHG and Zn-Al wire implants.

Surprisingly, the SHG material was the only material to elicit a relatively moderate amount of CD163 expression through the entire 6-month implantation time (Fig. 1-S1). The Al alloys all had consistently low expression of CD163 over the time course, with a large increase of expression over 6 months for the 4N material. The consistent presence of CD163+ cells around the SHG material could explain its positive tissue remodeling and integration, while the 4N response could be explained as an effort to reconFig. the arterial damage due to the necrotic response shown in Fig. 1-2.

Although cells had penetrated the corrosion layer at early time points in the SHG and Zn-Al materials, a decrease in cell viability can be seen from SHG, to Zn-1Al, and then Zn-3Al, respectively, with the SHG corrosion layer exhibiting the fewest irregular cells. The graded differences in cell viability within the corrosion layer of the Zn-Al alloys may explain the increasing capsule thickness from SHG and Zn-1Al to Zn-3Al, as well as the lack of a dense collagenous capsule for the SHG material, as seen in Fig. 1-3. Fig. 1-8 clearly shows high initial CD68 presence for the Al additions, which could be indicative of a highly aggressive macrophage attack in the early weeks of implantation. The graded difference in cell viability was associated with the degree of CD68 labeling, suggesting that Mφ activity may have contributed to the reduced cell viability within the corrosion layer. Although the presence of aluminum appeared to have elicited a greater inflammatory response, the Al³⁺ is not believed to be harmful, because there was no indication of toxicity to cells away from the regions of tissue where the macrophages had clustered. Potentially, the aluminum addition may have produced corrosion products (possibly Al₂O₃) that are too small or chemically stable for efficient phagocytosis, resulting in an increased residence time and build-up of Mφs in and around the corrosion layer. Another explanation is that the intergranular mode of corrosion may elicit a more robust inflammatory response relative to the surface corrosion experienced by the SHG material. For instance, oxygen gradients established by its transport into the metal interior through grain boundaries may stimulate secretion of harmful reactive oxygen

species by Mφs. Lastly, Zn-Al corrosion may increase the pH near the implant ²⁰, resulting in cell toxicity due to hydroxide ion transport from the material to the biological tissue.

1.5 Conclusions

Zinc implants produced dramatic differences in inflammation in rat arteries. The differences in biocompatibility of zinc materials are attributed in this study to their compositional dissimilarities that cause variations in implant corrosion activities and zinc ion fluxes. We conclude that the corrosion behavior (surface vs. intergranular), corrosion rate, corrosion product porosity, and the presence of a dynamic tissue-material interface regulate the inflammatory progression and remodeling of surrounding and penetrating tissue. Mφs and their monocyte precursor are known to orchestrate the fibrotic response in an up/down regulatory manner ²¹, making them the key cells that mediate inflammation and implant degradation. The ability of Mφs to penetrate and remain viable within the corrosion layer at early times may be of fundamental importance for eliciting long-term biocompatible inflammatory responses around corrodible metals. Future work will aim to further clarify the mechanisms by which zinc metallic implants regulate Mφ activity and viability with the aim to control inflammatory responses around vascular implants made from zinc base materials and predict long-term biocompatibility from early stage corrosion behavior.

1.6 Acknowledgements

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2 Development of novel in vivo methodology to screen degradable materials for biodegradable stent development^{3 4}

2.1 Introduction

Zinc alloy development and materials characterization for vascular stent application has been facilitated by a plethora of standardized and inexpensive methods. These methods have been employed over the past several years to generate a number of candidate zinc based alloys for vascular stents [1-3]. In contrast, the biocompatibility evaluation of zinc based degradable biomaterials for vascular applications has been challenged by a lack of standardized procedures at early developmental stages. While materials that have been selected for stent prototyping are commonly evaluated in a large animal stent implantation model using procedures designed for human subjects, such an approach is prohibitive for evaluating early stage materials due to its high cost and requirements of medical-grade facilities and medical expertise. With the steady accumulation of new zinc based materials [2], a standard and inexpensive approach for the biocompatibility testing of degradable metals for vascular applications is urgently needed.

Historically, the degradable metals community has relied upon in vitro evaluation of candidate materials to assess biocompatibility. In this fashion, cell culture approaches dominate the preliminary biological testing, with investigators focusing their attention on either cytotoxic or mechanistic investigations. Cultured cells are exposed to Zn^{2+} ions [4-6] or corrosion extracts collected and concentrated from candidate materials that were corroded in one of a number of different corrosion solutions [7-10]. These have become standard approaches that can provide fundamental insight into the interactions between corrosion products and various cell types.

³ This chapter contains material previously published in the Journal of Metals

⁴ References in this section follow the author guidelines for JOM

While directly investigating cell responses to corrosion products provides a fundamental understanding of the biologic response, results are not directly applicable to the compatibility of the material in the application environment. An artery is a complex tissue with three main resident cell types (media - vascular smooth muscle cells, intima - endothelial cells, and adventitia - fibroblasts). While the cytotoxicity and sub-cellular response of any of these three cell types is important for determining the overall compatibility of the tissue to degrading zinc materials, molecular cross talk, differences in proximity between the three arterial compartments to the implant, and interaction of blood borne cells play an important role in macro-level vascular responses, which cannot be replicated in vitro [11]. For instance, activation of an organism's innate immune system by biomaterial implantation and tissue injury will impact the overall reaction against the material, critically affecting device performance [12]. Furthermore, the corrosion of zinc based materials varies widely between corrosion solutions and physiological conditions. A distinction must be drawn between cell toxicity (obtainable from direct exposure of cultured cells to Zn^{2+} or corrosion products) and application specific biocompatibility (requiring in vivo assessment of the material in the appropriate physiological environment).

Stents implanted into the arterial environment are expected to restore and maintain continued function of the artery. Harmful events, including smooth muscle cell neointimal proliferation, thrombogenesis, negative arterial remodeling, and destructive inflammation contribute to the narrowing of the lumen and restenosis of the stented segment. While measurements of cytotoxicity can contribute to predicting negative stent outcomes, it cannot replace direct application-specific testing. For instance, magnesium and iron biometals perform superbly when measured for cytotoxicity, but struggle in the application environment due to drawbacks in their corrosion rates that eventually lead to device failure [13, 14].

In 2012, our group introduced a metal wire implantation model as a cost effective and simplified approach for the biocompatibility evaluation of degradable materials in the vascular environment [15]. Any material that can be drawn into a 0.20-0.25 mm nominal

diameter wire can be evaluated in this model to investigate material biocompatibility. Implanting a wire into the lumen of the abdominal rat aorta rather than a stent into a pig is promising from a device testing standpoint: the wire affords a simpler geometry without balloon injury, and is less expensive in terms of animal costs, materials preparation, and corrosion analysis.

In this contribution, we aim to describe a cost effective histomorphometric standard for a pipeline-based evaluation of candidate degradable materials using the murine based wire implant model. Herein, wires made from two zinc alloys and pure zinc were implanted into the abdominal aorta of Sprague Dawley rats and collected at 6 and 11 months. Neointima tissue surrounding the wire was measured for area, thickness, and protrusion into the arterial lumen, critical metrics for quantifying vascular application based biocompatibility.

2.2 Experimental

2.2.1 Materials

Staining reagents include powdered iron hematoxylin (H3136), sodium sulfate pentahydrate ($\text{Na}_2\text{SO}_4 \cdot 5\text{H}_2\text{O}$) (753599) anhydrous iron (III) chloride (FeCl_3) powder 99.99% (157740), phosphate buffered saline pH 7.4 (PBS, P4417) , Lugol's iodine solution(624-71), xylene substitute (78475), Van Gieson solution (HT254), and Eukitt quick hardening mounting medium (03989); all were obtained from Sigma Aldrich(Burlington, MA). 200 proof Ethyl alcohol (absolute ethanol) was obtained from Pharmco-Aaper. Neutral buffered formalin 10% (HT501128) was also obtained from Sigma Aldrich.

2.2.2 Surgical Implantation

For the present study, a complete histological characterization of metallic wires implanted into the rat abdominal aorta was explored. Pure zinc (Zn) (number of samples) =10, zinc-4 wt. % lithium (Zn-4Li) (number of samples) =10, and micro-alloyed zinc-

magnesium (number of samples) =12 (either Zn-0.01 wt. % Mg or Zn-0.1 wt. % Mg, hereafter referred to as Zn-XMg) wires were implanted and collected at 6 and 11 months. Two sample sites were evaluated per wire (near both puncture sites for each wire) resulting in n=64. Greater details of the Zn, Zn-4Li, and Zn-XMg wires used in the present study can be found in references [16, 17].

Fig.2-1 Panels 1-9 show the surgical depiction of wire implantation. Yellow “V” and blue “A” denotes the inferior vena cava and descending abdominal aorta respectively. Purple “W” highlights a wire for insertion. Panel A shows the Sprague Dawley rat in the supine position with the abdominal area shaved and

cleaned before procedure. B shows the abdominal muscle with multiple sutures. C shows the skin clamped with stainless steel staples.

All animal experiments were approved by the Michigan Technological University IUCAC board. Rats were anesthetized in the supine position using an O₂ rich isoflurane vaporized gas mixture. The abdomen was shaved and cleaned using antiseptic wipes (Fig.2-1a). For the initial incision, the skin was cut with dissecting scissors through the midline starting approximately 3 cm above the pelvic region and terminating after approximately 5-8 cm. The abdominal muscle was cut in a similar fashion (step 1 of Fig.1). With the viscera exposed, intestines were carefully pushed aside in order to visualize the caudal descending abdominal aorta and vena cava (step 2). Connective tissue surrounding the vessel was carefully pushed aside with Q-tips to completely expose the vessel. In order to continue with the implantation, blood flow must be briefly interrupted, which is accomplished by placing vascular clamps inferior to the renal arteries and superior to the aortic bifurcation. First, the aorta and vena cava must be carefully separated by using blunted forceps. This is accomplished by carefully poking through and then enlarging an opening between the vessels to fit the clamp. After separation, the proximal clamp is placed first, then the distal clamp applied (steps 3-6). A wire sharpened by angular cuts (using very sharp scissors) is poked into the artery at a location superior to the distal clamp (steps 7 & 8). After insertion, the wire is advanced within the artery and then punctured out of the artery, producing a second puncture site. The wire is left in place with ~2 - 3mm of wire exteriorized from the artery at both ends. Zinc wires do not need to be secured in place, because corrosion activity embeds the wire within the tissue. If biostable wires are implanted (as controls, for example), the wires must be secured in place to prevent shifting. The vascular clamps are removed (the distal clamp is removed first). The abdominal muscle is sutured together, and the skin clamped (panels b & c). The rat is allowed to recover under observation, following a subcutaneous injection of butorphanol for pain management. Proper mobility of the lower limbs is used as an indicator of continued abdominal aorta function. All animals survived the procedure.

2.2.3 Implant removal and gross examination

Rats are anesthetized using a vaporized isoflurane and O₂ rich mixture. After the subject is deeply anesthetized, the abdominal aorta is isolated from the surrounding tissue. The diaphragm is punctured and the heart is removed from the thoracic cavity to ensure death. The aorta containing the wire is then collected and examined grossly for any obvious abnormalities. The aorta is then placed in a small cryo-mold, surrounded by optimum cryo-temperature cutting medium, snap frozen in liquid nitrogen, and stored at -80°C until cryo-sectioning.

2.2.4 Tissue preparation and histological evaluation

Specimens that are snap-frozen can be stored at -80°C for up to one year while maintaining tissue integrity. To prepare for histological evaluation, the implant containing aorta is oriented to ensure a cross-sectional profile of the wire/artery. The artery is then freeze-mounted on a specimen chuck, and placed onto the head of a cryo-microtome for cryo-sectioning. In our experience, a blade angle of 12 degrees and a chamber temperature of -24°C to -26°C is optimal for these samples. A section thickness of 5-10 µm can be reliably produced, but for the present study all sections were taken at a section thickness $\approx$ 8 µm. We initially section rapidly through the exteriorized portion of the wire while visualizing the wire in the sample (with a hand-held magnifying lens) as it moves progressively closer to the artery. Once the wire in the specimen can be seen within the wall of the artery, the sample is more carefully sectioned until the wire appears to pass the internal elastic lamina. The wire position in the sample can be verified by inspecting cross sections on an inverted microscope equipped with polarized light filters. When the wire has crossed the internal elastic lamina, sections are collected on slides for later staining, with multiple sections per sample collected.

Fig.2-2 VVG staining, Panels 1-3 highlight the sectioning progression of the wire position in relation to the artery. Red “w” shows the Zn wire and yellow arrows are positioned at the first elastic fiber. Cartoon depiction displays four most common neointima types, with the explained morphometric measurement locations in relation to the neointimal formation.

This process is seen in Fig.2-2, with clear progression of the wire into the luminal space of the artery. To ensure a representative characterization is performed, cross sections are collected over a distance of 100-500 μ m into the specimen. As the sample is

progressively cross-sectioned, the wire may eventually appear to “free float” in the arterial lumen. Cross sections where the wire is not in contact with the artery are not used for histological analysis. Cross sections near the second puncture site can be collected either by flipping the specimen over on the chuck and repeating the procedure described above, or by continuing to section through the sample, although this is more labor intensive. By this method, cross sections near both puncture sites from the implant can be obtained.

2.2.4.1 *Histological Staining and Morphometric Analysis*

Slides with frozen sections were fixed for 10 min at room temperature using a buffered-formalin solution. After fixation, the slides were washed 3 times for 5 min each in phosphate buffered saline. A hematoxylin and eosin stain (H&E) was carried out as a general stain in order to visualize tissue constituents and qualitatively assess vessel morphology (see the following reference for procedure details [18]). A Veorohoff Van Geison elastic tissue stain (VVG) was carried out to identify the elastic laminae within the artery. This provides anatomical landmarks for the histomorphometric analysis.

For Veorohoff Van Geison staining, slides with tissue sections were fixed in 10% neutral buffered formalin for 10 minutes. Slides were then washed in three changes of PBS for 5 min each and overstained in Verhoeff’s elastic stain for 10 min (30 mL of absolute ethanol and hematoxylin solution 5% m/v, 12 mL of deionized water and FeCl_3 solution 10% m/v, and 12 mL of Lugol’s solution). The slides were then differentiated using an inverted optical microscope with a FeCl_3 solution diluted to 2% m/v until elastic fibers in the media layer of the arterial cross-section was well defined, and non-elastic tissue became unstained. The slides were then rinsed in deionized water for 5 min. The slides were placed in a deionized water and $\text{Na}_2\text{SO}_4 \cdot 5\text{H}_2\text{O}$ solution, 5% m/v for 1 minute and then rinsed in running tap water for 5 min in order to “blue” the hematoxylin. The slides were counter stained in Van Gieson solution for 3 m and microscopically differentiated in a 95% v/v ethanol solution until there was a clear distinction between the pink Van Gieson stained adventitia and the brown stained media. The slides were dehydrated in

two changes of absolute ethanol for five min each. The slides were then optically cleared with two changes of xylene substitute for 5 min each and mounted under a coverslip using Eukitt quick hardening mounting medium. The samples were imaged using an Olympus BX51, DP70 bright-field microscope (Upper Saucon Township, PA).

Neointimal area (NA), wire-to-lumen thickness (WLT), and base neointimal length (BNL) were measured using METAMORPH software. Cross-sections stained with VVG and imaged at 100X normal magnification were used for the measurements. The neointima was considered as tissue surrounding the implant on the luminal side of the first elastic fiber, which was highlighted by VVG staining, with the wire cross sectional area excluded. All neo-tissue growth contacting the implant was included in the measurement, as well as all significant intimal activation in nonadjacent regions. WLT was measured near the apex of the neointima, at the point where the tissue protruded furthest into the lumen. For BNL, the first elastic fiber was traced (below the wire implant) until negligible neointimal activation was observed (Fig.2-2). Multiple sections of each sample were evaluated to ensure a representative depiction of each sample (minimum of 5 sections). Only sections that satisfied quality concerns (section not torn considerably, neointima still intact, low amount of section folding) were measured.

2.2.4.2 Zn Accumulation in Organs

To provide insight into whether zinc is cleared or retained with the body during implant degradation, the organs that play a major role in the distribution, filtering and regulation of the fluid compartment and constituents were analyzed for zinc concentration. Wires were implanted within the abdominal aorta for 6 months, whereupon the heart, blood plasma, liver and kidney were collected. Using a fluorometric zinc quantification assay (ab176725) and a Bradford assay (ab102535), all purchased from Abcam (Cambridge, MA), zinc levels were determined relative to control tissue (collected from non-implant rats), and reported in $\mu\text{g/mL}$.

2.2.5 Statistical Analysis

Histomorphometric data was uploaded into MATLAB R2018a. A Wilcoxon ranksum test was performed on all test groups within each respective time point (6 or 11 month implantation) relative to the reference Zn implant. A p value <0.05 was considered significant. Data plotted in boxplot graphs show the upper and lower quartiles (25%-75%) within the boxed region, 0% and 100% quantiles within the whiskers, with red '+' as computed outliers. For tissue derived Zn^{2+} organ concentration, a two-tailed student's t-test with assumed equal variances was used to determine significance between control and test groups. Bar graphs denote sample mean, with error bars in standard deviation. A p value <0.05 was considered significant.

2.3 Results and discussion

The development of zinc-based metals is advancing in academic institutions although it has not yet reached the clinical stage. The potential for targeting multiple vascular applications as degradable vascular implant materials has sparked intense interest amongst metallurgists and corrosion scientists with expertise in materials development. To the authors' knowledge, most of the in vivo work published to evaluate zinc based materials in the vascular environment has emerged from our group, with a single contribution based on pure Zn stents deployed in the rabbit abdominal aorta by Yang et al, [19]. In order to develop a coherent framework for the early in vivo testing of zinc implant materials, we have provided extensive details of our small animal model, including surgical, tissue sectioning, and histological staining procedures. We have also developed metrics to quantitatively characterize the long-term histological progression of the neointima that forms around zinc based wire implants. The approach used here could become a standard in the field, which would allow direct comparison between biodegradable materials developed by different laboratories throughout the world. The metrics we have developed for the wire model have been adapted from metrics that are used to characterize the neointima that forms around vascular stents. Therefore, these

metrics may help predict performance of the same material when deployed as a stent in a large animal model.

2.3.1.1 *General histological presentation*

The evaluation of neointimal characteristics provides critical insight into the biological performance of intraluminal implants. Stent performance is routinely evaluated by specific morphometric characteristics including lumen area, neointimal thickness, neointima area, and internal elastic lamina area [20-22]. The wire implantation model simulates a single stent strut placed within the lumen of an artery, without concomitant radial luminal injury or mechanical excitation of the implant. As it is known that radial force exerted during balloon catheter deployment provokes inflammation and smooth muscle cell proliferation[23], the present model allows us to investigate the host response to the experimental material without confounding effects related to deployment. Classical stent metrics such as lumen area and external elastic lamina area (used to determine positive and negative remodeling) are not useful when using the wire model, due to the wire not engaging the entire artery circumferentially. The characteristics of the developed neointima surrounding the wire implant provides the most useful information relevant to implant performance. Fig.2-2 (lower box) summarizes the various neointima types that dominate most observations in our wire model. Type 1A neointima formations are considered ideal, with a low profile and a stable, mature tissue response to the implanted wire. Type 1B formations are common at longer time points (11 months), and in our experience are believed to be the natural temporal progression of Type 1A formations. Type II formations reflect the negative responses seen in this model. The characteristics of Type 2A are assumed to be analogous to negative arterial remodeling of a stented artery, which contributes to luminal area reduction and reduced vascularization. We have seen this response in our previous investigation of Fe wires and in a more dramatic fashion with PLLA coated Zn wires [15, 24]. The failure mode of this formation consists of vascular smooth muscle cell (VSMC) neointima hyperplasia, which we confirmed by alpha actin positive staining (data not published). Type 2B presents as a thin neointima with voluminous corrosion product and a cell rich middle layer. The consistent thin tissue

surrounding the implant could result in rupture, which is not ideal for stenting applications.

Fig.2-3 H&E staining, 4 representative sample cross-sections of each implant at 6 and 11 month residence. Images are taken at 100X normal magnification and scale bar is approximately 1mm.

In order to properly characterize neointima formations, thin tissue sections must be prepared and stained for each sample. Hematoxylin and eosin is the most common, general histological stain and we have routinely used this stain to evaluate materials implanted in the rat model [15-18, 24, 25]. Fig.3 shows multiple representative cross sections of 6 and 11 month samples that have been stained with H&E in order to visualize neointimal formations. Inflammation and macro-level cellular responses can be readily identified at high magnification in these H&E images, which we have demonstrated elsewhere [18, 25]. Considering the presence and distribution of cellular type and inflammatory responses directly influence neointimal formations and contribute to the compatibility of intraluminal implants [23], a reliable and straightforward workflow of neointimal morphometric characteristics can be used to describe the overall cellular response to the material. To make measurements that describe the neointimal character, a VVG stain is more appropriate because this stain can be used to distinguish the different layers of arterial tissue.

Fig.2-4 VVG staining, 1 representative sample at each time point for the alloys investigated. "W" denotes wire position. Scale bar is approximately 1mm

The elastic fibers (black) provide anatomical landmarks for NA and BNL quantifications, which are also identifiable but far less prominent on an H&E stain. Fig.4 shows a representative VVG stain of each implanted alloy at 6 and 11 month time points. A general progression of Type 1A to 1B formation can be seen for the Zn wire. The Zn-4Li specimens generally display increased intimal activation away from the implant, with a reduction of elastic fibers near the implant. For Zn-XMg wires, large Type 2A and 2B formations are often seen at both time points, with a high variation in neointimal responses.

Fig.2-5 VVG staining of samples at indicated lumen obstruction index (LOI). Failure threshold is shown at 30%. Tabulated morphometrics for the three alloy systems investigate at 6 and 11 months.

2.3.1.2 Identification of optimal performance

In order to quantify the direct success or failure of candidate materials, a lumen obstruction index was developed to describe negative histomorphometric appearances.

To obtain the lumen obstruction index of a sample:

$$\text{LOI} = \left[\frac{\text{WLT}}{\text{BNL} - \Phi} \right] \times 100\% \quad (1)$$

where WLT is the wire to lumen thickness, BNL is the base neointimal length, and Φ is the nominal wire diameter, here as 250 μm . Presumably, a native artery that lacks a wire implant and has normal intima anatomy would have an LOI of 0%. An index of 30% was set as the cutoff for degradable material success. This cutoff was based on the largest index measured for a pure Zn wire implant, out of 60 observations. Visual depiction of the failure index is presented in Fig.2-5, with the cutoff (dashed red line) identifying the failed pure Zn neointima. Out of all observations at both time points, one Zn implant at 6 months and three Zn-XMg implants at 6 months exceeded the failure threshold. These implants were not included in the subsequent statistical analyses.

Fig.6 shows measurements from histological sections for samples at the two long-term evaluation time-points. At 6 months, no statistical differences exist between the Zn reference and Zn-4Li / Zn-XMg alloys for any metric. Additionally, the average WLT for Zn and the Zn-4Li/Zn-XMg alloys appears modest ($67 \pm 36 \mu\text{m}$ vs $47 \pm 28 \mu\text{m}$ / $42 \pm 14 \mu\text{m}$ respectively). All three failed Zn-XMg implants and the one failed Zn implant occurred at 6 months, indicating this time point could be useful as the minimum to observe failure. For the 11-month implants, all metrics for the alloys were significantly increased relative to the pure Zn reference material. WLT for the Zn-4Li and Zn-XMg systems ($90 \pm 59 \mu\text{m}$ and $77 \pm 38 \mu\text{m}$) is significantly higher than for Zn, $60 \pm 69 \mu\text{m}$ at

Fig.2-6 Boxplot representation of metrics at each time point, for each alloy system. The upper and lower quartiles (25%-75%) are within the boxed region, 0% and 100% quantiles within the whiskers, with red '+' as computed outliers. Significance was determined by a Wilcoxon rank sum test, with p values in respect to the reference zinc sample at each time point. "*" $p < 0.05$, "***" $p < 0.005$, "****" $p < 0.0005$.

11 months. Zn WLT appears unchanged from 6 to 11 months, while this metric appears to have increased markedly for Zn-4Li and Zn-XMg implants.

Neointima thickness is commonly used to gauge arterial responses to stenting due to its correlation with stent failure and luminal area reduction [26-28]. In the present study, neointima thickness progression measured by WLT appears to be a reliable metric in

determining compatibility. Furthermore, when divided by the length of neointimal activation (BNL) it can serve as an indicator of negative remodeling (closer to 100% indicates excessive protrusion within the lumen). The area of the neointima (NA) is important, but does not provide enough information to adequately describe performance of the material. For instance, two implants can have similar NA measurements but dramatically different failure indices. Therefore, NA is not recommended as a stand-alone metric for compatibility, but as a complementary evaluation.

Although a failure threshold of 30% is recommended, neointimal formation values greater than 20% should denote poor performance. Although none of the Zn-4Li implants met the failure threshold, multiple instances of increased WLT with failure indexes close to 20% (data not shown) suggest a reduction in compatibility for the Zn-4Li wires relative to pure Zn.

Fig.2-7 Zn Organ quantification by a flourometric assay. Bar graphs represent mean and error bars are $\pm$ standard deviation of the samples. Minimum n=3 for each condition, significance was determined by a two tailed students t test, “NS” not significant

An organ analysis can also be performed on each animal, as shown in Fig.2-7. This analysis demonstrated a slight increase in plasma bearing Zn, with no accumulation of zinc in the heart, liver, or kidney. This finding suggests that zinc removed from the implant is safely transported and excreted from the body.

Fig.2-8 Workflow of wire model showing alloy development and biocompatibility testing

The wide variety of Zn-alloy systems that can serve in degradable cardiovascular stent applications has opened exciting possibilities for a number of candidate materials. The most appropriate approach for assessing the biocompatibility of degradable metals remains an important and open question for materials selection. Here, we present a robust workflow, summarized in Fig.2-8, for the testing of degradable metal candidates. Candidate materials are designed, extruded into rods, drawn to $\phi=0.25$ mm wire, and characterized for mechanical and microstructure properties. The wires are then implanted into the lumen of Sprague Dawley rats and allowed to maintain residence for up to approximately 1 – 2 years. Implants are then collected and histologically processed for biocompatibility assessment. Based upon quantification of metrics as outlined in the present contribution, informed decisions related to alloy design are iterated into the workflow, or selected materials can progress to stent testing in large animals. A summary of these proposed metric values are given below.

Table 2-1: Summary of the proposed biocompatibility values given by the wire model

Metrics	Biocompatibility standards (6-months)
Lumen obstruction index (LOI)	$\leq 30\%$
Wire to lumen thickness (WLT)	$\approx 70 \mu\text{m}$
Neointima Type	Type 1 (A or B)

Based upon the metrics developed here, pure Zn elicited the best biocompatibility of the three evaluated materials. An excellent biocompatibility response for pure Zn in the wire model is consistent with what was reported for pure zinc stents implanted within the

rabbit abdominal aorta [19]. Therefore, the metrics we have developed based on implant materials in wire geometry may be predictive for the biocompatibility of stents deployed in large animals. Zn-4Li exhibited a reduction in biocompatibility at 11 months relative to pure Zn due to the progression of its WLT. Although no Zn-4Li wires incurred a LOI > 30%, numerous implants incurred elevated LOI measurements above 10%. The present finding of a non-obstructive neointima is consistent with previous preliminary observations of Zn-4Li wires made by our group [17].

Zn-XMg performed the worst of the three implanted materials, based on three failures at 6 months, as determined by LOI, and a significant increase of WLT at 11 months relative to pure Zn. This reduction in compatibility was also reported previously, and may be due to an increase of inflammation in Zn-XMg implants [16]. Due to the well-known compatibility of Mg, the authors speculate that the performance reduction was unrelated to the inclusion of Mg directly. Rather, the worsened response may have been caused by changes in corrosion behavior or the generation of intermetallics of reduced biocompatibility. These factors gave rise to Type 2A and 2B neointima formations, even within the same sample (not shown), confirming our earlier work highlighting the importance of corrosion behavior for regulating biological responses[18].

2.4 Conclusions

We conclude that:

Biocompatibility can be reliably determined using the rodent wire implant model

Six months is the minimum viable time point to confidently determine biocompatibility

Quantifications of neointima characteristics, including wire to lumen thickness and base neointima length, can describe the overall performance of degradable implants.

A reduction in biocompatibility relative to pure zinc is seen for the Li and Mg alloy systems.

2.5 Acknowledgments

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3 Contributions of the surface characteristics of degradable zinc implants to the neointimal response and subsequent biocompatibility^{5 6}

3.1 Introduction

Zinc (Zn)-based metals are emerging materials for biodegradable cardiovascular devices¹⁻⁴. Although many questions remain, a number of recent preclinical studies have reported an encouraging efficacy for pure zinc and Zn alloy based stents⁵⁻⁷. Additionally, Zn-based metallic implants are steadily closing in on the mechanical benchmarks required for stenting human arteries⁷⁻⁹. However, in addition to mechanical strength and ductility, degradable Zn-based stents may require an engineered surface oxide film to initiate a relatively uniform corrosion and biocompatible host response in early stages of implant-tissue interactions¹⁰⁻¹². While there has been considerable progress in the development of novel Zn-based degradable alloys with improved mechanical and structural properties, the development of surface films has consistently lagged behind^{10, 13}. This is probably due to widely accepted standards for permanent stents related to smooth and electropolished surfaces, which may not apply directly to degradable metals.

A key feature of degradable Zn is an in vivo corrosion rate that is sufficient for device dissolution within 1-2 years¹¹. Interaction with the host occurs at the Zn implant-tissue interface throughout the service lifetime, although the Zn-based substrate interface changes over time¹⁴. Continuous corrosion at the implant interface and the production of various biodegradation products elicit a host response that is sustained until the implant is fully degraded. It is well known that a change in the passivity or susceptibility to corrosion is determined largely by the surface film characteristics¹⁵⁻¹⁷. However, the ideal surface properties for degradable Zn materials remain an open question. Critically, a

⁵ This chapter contains material previously published in ACS Applied Materials and Interfaces

⁶ References in this chapter follow ACS author guidelines

surface that inadvertently provokes a rapid or non-uniform corrosion of the underlying degradable Zn substrate could result in localized metal dissolution and changes in pH. These processes, if not controlled, may be harmful to surrounding cells, initiate inflammation and/or neointimal growth, and increase susceptibility of implant to stress corrosion cracking. In contrast, an improved biological response and mechanical performance could be imparted from engineered Zn surfaces that increase corrosion resistance and corrosion uniformity.

Conventional biostable stents are commonly surface treated by electropolishing (EP) to create a smooth surface finish with a thin, stable oxide layer that protects the underlying metal from corrosive attack¹⁸. Since degradable metals are designed to corrode, a thin EP surface layer may be highly sensitive to local variations in cells and tissue composition. In contrast to EP, the anodization (AD) of metal surfaces may increase the stability of the surface layer and therefore reduce its sensitivity to non-uniform penetration from local variations in the biological milieu^{19, 20}. Both EP and AD processes are widely used in the biomedical industry as surface finishing treatments^{10, 21}. During EP, the surface material is removed by anodic dissolution leaving a smoother surface with reduced roughness and improved cleanability²². AD consists of an electrolytic passivation, which is used to increase the thickness of the natural oxide layer of the implant. Both EP and AD procedures have been reported to dramatically improve the biological performance of permanent²³⁻²⁶ and degradable implants.

The EP and AD protocols are well developed for inert implant materials such as stainless steel, titanium alloys and cobalt alloys. However, they are in their infancy for Zn-based implant materials. Here, we describe methods to exert control over the formation and uniformity of surface oxide films on Zn material surfaces using electropolishing and anodization processes. Zinc materials with engineered surfaces were characterized by electrochemical methods. They were then implanted into Sprague Dawley rats for 2-8 weeks in an effort to correlate neointimal responses to surface oxide film characteristics. We found that anodization of Zn implants, although not commonly applied to vascular stents, could benefit the biocompatibility of Zn-based stents if engineered appropriately.

3.2 Experimental

3.2.1 Surface preparation

High purity (99.99%) Zn wires ($d = 0.25$ mm) and discs ($d = 8$ mm), ethanol (C_2H_5OH), butanol (C_4H_9OH), aluminum chloride hexahydrate ($AlCl_3 \cdot 6H_2O$), zinc chloride ($ZnCl_2$), oxalic acid ($(COOH)_2$), and industrial detergent (Citranox[®]) were purchased from Sigma Aldrich (St. Louis, MO, USA). The EP process was optimized bearing in mind criteria that provide reduced surface roughness and dimensional precision aimed to minimize thickness loss, which is of paramount importance for thin wires and stents. The electrolyte was designed based on the indications of ASTM E1558-09 standard²⁷. Regarding anodization, the process parameters were selected to allow maximum surface coverage with minimum coating thickness. The study consisted of Zn wires and discs treated under a combination of different voltages, mixing speed, electrolyte concentration and treatment duration. The EP and AD sample surface finish was assessed by microscopic observations. Subsequently, the samples exhibiting desired surface properties were selected for in vitro and in vivo degradation study. Zn samples were cleaned with a detergent, then subjected to an ultrasonic cleaning process in pure acetone, followed by rinsing with deionized water. After that, the specimens were electropolished with an electrolytic solution comprised of C_2H_5OH (885 mL), C_4H_9OH (100 mL), $AlCl_3 \cdot 6H_2O$ (109g), $ZnCl_2$ (250g), and distilled water (120mL). A 20 x 20 mm 316L stainless steel sheet was used as a cathode. The distance between the electrodes was ~60 mm. The current was 0.45 A, corresponding to a voltage of ~26–28 V, and the temperature was ~21–23 °C. The duration of the procedure was ~90 s. Throughout the course of electropolishing the solution was stirred with magnetic agitation to avoid waviness and streaks on the surface. Prior to anodization, Zn discs and wires were electropolished. The samples were submerged in 1L of 0.5 M $(COOH)_2$. $(COOH)_2$ was selected as an electrolyte because of the oxalate ability to bind strongly to the metallic surface, thus improving the substrate corrosion resistance²⁸. Anodization occurred for 1 min at room temperature using a 316L stainless steel cathode. The power supply was set to a current of ~4 A corresponding to a voltage of ~10V. During anodization, the

electrolyte solution was constantly stirred with magnetic agitation to reduce the thickness of the double layer at the metal–electrode interface to obtain uniform local current densities on the Zn electrode. All samples were cleaned in an ultrasonically stirred acetone bath for 3 min and in a 75% (v/v) ethanol bath for 5 min after each stage of preparation.

3.2.2 Surface characterization

A FEI Philips XL 40 equipped with a tungsten filament and operated with an acceleration voltage of 15 kV was used for scanning electron microscopy (SEM) analyses.

Micrographs were acquired with a probe current in the range of $1 \cdot 10^{-10}$ to $1 \cdot 10^{-8}$ mA. X-ray photoelectron spectroscopy (XPS) characterization was carried out to determine the chemical composition of the EP and AD surfaces (PHI 5800 spectrometer, Physical Electronics U.S.A), with an incident angle of 45° with respect to the surface, and a residual pressure of $5 \cdot 10^{-9}$ Torr. A survey spectrum was recorded using a standard magnesium X-ray source for survey spectra (0–1400 eV) at 400 W to identify all elements present at the surface. High resolution spectra (HR) of C1s, O1s and Zn p_{3/2} regions were recorded with a standard Mg K α X-ray source (1253.6 eV). The spectrometer work function was adjusted to give 284.8 eV for the main C1s peak. Curve fittings were determined using the Gaussian–Lorentzian (80–20) function and a Shirley type background and performed using the software Multipack[®]. Each peak was decomposed into Gaussian/Lorentzian components, which have been attributed to the presence of oxides, hydroxides, and carbonates/water. C 1s peak fitting was carried out with following components: peak I C-C/C-H (~284.8 eV); peak II: -O-C=O- (~289.0 eV), and peak III: carbonate groups and/or adsorbed CO and CO₂ (~290.1 eV²⁹⁻³¹). The oxygen peak was composed of contributions located at: peak I: ~530 eV (oxide); peak II ~531.5 eV (hydroxide); peak III: ~533.5 eV (carboxyl groups, carbon dioxide and/or adsorbed H₂O^{31, 32}). For Zn 2p_{3/2}, the subsequent peaks were identified: peak I (~1021.8) metallic zinc, peak II (~1022.) zinc oxide and peak III (~1022.7) zinc hydroxide and/or carbonate.

3.2.3 In vitro degradation behavior

The corrosion of EP and AD samples was studied in Hanks' solution supplemented with 0.35g NaHCO₃/L. The pH of the solution was adjusted to 7.4 using 1 M NaOH or HCl. Open circuit potential (OCP), electrochemical impedance spectroscopy (EIS), and potentiodynamic polarization tests were performed to study the corrosion behavior of the investigated samples. The potentiodynamic test was performed using a conventional three-electrode cell (Princeton Applied Research Model K47) with a platinum counter electrode of 1 cm² in surface area, a saturated calomel reference electrode and the prepared Zn-based working electrode. The OCP vs the SCE RE was monitored without applying any outside source for 3600s until equilibrium was reached at the corrosion potential E_{corr} . A scan rate of 0.166 mV/s, with an applied potential range of 1V, was used. The experiments were carried in an aerated environment at 37±1°C. The solution was stirred with magnetic agitation (80 rounds/min) during the test. For each type of material, three specimens were tested using the same conditions. The electrodes were connected to a potentiostat (PARSAT4000, Princeton Applied Research, PA, USA) and monitored using the VersaStudio® software. The corrosion rates were obtained based on the calculated corrosion current density (i_{corr}), using the following equation:

$$CR = 3.27 \cdot 10^{-3} \frac{i_{corr} EW}{\rho} \quad (\text{Eq. 1})$$

where CR is the corrosion rate (mm year⁻¹), i_{corr} is the corrosion current density (μA cm⁻²) obtained on the base of potentiodynamic curves using Tafel extrapolation method, EW is the weight equivalent and ρ is the material density (g cm⁻³).

Impedance spectra were separately collected from 100 kHz to 0.1 Hz with 5 mV perturbation amplitude. Because of reactive nature of zinc, no data points were collected below 0.1 Hz. The OCP potential was recorded for 30 min prior to the EIS measurement to allow the interface to reach a steady state. EIS spectra were recorded after 1h, 24h, 72h, 120h and 168h. The results were adjusted according to equivalent electric circuits using the software ZSimpWin®.

For static immersion tests, the Zn-based discs with 0.8 cm² exposed surface area were immersed for 28 days in 40 mL of Hanks' solution. The whole volume of solution was changed every 7 days to keep the pH value close to 7.4 and to maintain conditions as constant as possible. The containers were stored in a controlled temperature ($T = 37 \pm 1^\circ\text{C}$). Three specimens were tested for each condition. The samples were subsequently washed with 200 g L⁻¹ of chromium oxide in distilled water to remove corrosion products. The surface morphologies and chemical compositions before corrosion products removal were examined by SEM and energy dispersion spectrometry (EDS).

3.2.4 In vivo wire implantation

Adult female Sprague Dawley rats were purchased from Harlan Laboratories. Nine samples per treatment group were harvested over 2, 4, and 8 weeks (3 samples at each time point), using a surgical procedure described previously³³. At the euthanization time point, the aorta containing the wire was dissected and excised, surrounded in polyfreeze cryo medium, snap frozen in liquid nitrogen, and stored in a -80 centigrade freezer until cryo-sectioning.

3.2.5 Tissue preparation, histological, and morphometric analysis

200x magnified H&E images were uploaded to MetaMorph image analysis software. The thickest region of the neointimal growth was measured and recorded from cross sections taken from at least three different locations along the wire, spanning approximately 0.2 mm. A detailed description of the methods for the morphometric analysis are described in our previous publication³³.

3.2.6 Statistical Analysis

All data was uploaded into MATLAB with a statistics machine learning toolbox package. A Wilcoxon rank sum test was used to compare experimental groups. The lumen occlusion index was used as a cutoff to exclude outliers ($\text{LOI} \geq 30\%$)³³.

3.3 Results

3.3.1 Surface characterization

Fig.3-1. SEM micrographs of (a) EP and (b) AD samples, (c) Chemical composition of EP and AD Zn surfaces by XPS and overlay of XPS high resolution spectra and peak positions for (d) carbon – C1s, (e) oxygen (O1s), (f) zinc (Zn 2p_{3/2}) and (g) zinc auger (ZnLMM), (h) HR-XPS calculated values from the Gauss–Lorentz fit for the O1s and C1s peaks.

SEM micrographs of the EP and AD surfaces are shown in Fig. 3-1. A smooth surface, similar to the industry standard of bare metal biostable stents, is evident on degradable Zn due to the EP process. In contrast, the Zn AD surface (Fig. 3-1b) is composed mainly of prismatic particles (with sizes in the range of ~500-800 nm), similar in morphology to what has been produced by other researchers following Zn anodization in oxalic acid^{34, 35}. An ultra-thin oxide film (~4 nm) is formed on the EP surface (Fig. 3-2a), estimated using an XPS depth profile. The AD coating is relatively uniform, ~6 µm thick, and indicated by green two-headed arrow in Fig. 3-2c.

According to the XPS survey spectra, shown in Fig. 3-1c, Zn is present on both EP and AD surfaces. Non-metallic elements such as C, N, and Cl were detected on the surface of the both samples. Their presence is most likely a consequence of the surface preparation, as EP and AD treatments involve the use of different acids and solvents. Fig. 3-1d-g show the high-resolution scan data for C 1s, O 1s, Zn 2p_{3/2}, and Zn LMM peaks. The amount of oxygen on the surface is significantly higher for the AD sample, implying a higher content of oxygen-containing compounds. Contamination from hydrocarbon compounds was observed on both surfaces. Its source is constituents of the ambient air. The presence of carbon has been observed on clean surfaces of most materials, even under moderately high vacuum conditions³⁶. These carbon species are not involved in any chemical interaction with the underlying virgin surface species, as they are not very reactive by their inherent chemical nature³⁷. The content of carbon was lower for AD samples as seen in Fig. 3-1c. In contrast, the air exposure of the EP surface resulted in rapid accumulation of C containing compounds, because chemically evolving clean metallic surfaces are more vulnerable to opportunistic carbon contamination³⁷. The C1s

HR spectra for EP samples reveal the main component as a C-C/C-H peak at 284.8 eV. The AD surface treatment resulted in a carbon presence, with the main peak appearing at a binding energy of 289.4 eV. This peak can be divided into two components: C in the carboxyl group (-O-C=O-) ascribed to Zn oxalate and/or carbonate groups and carbon at 290.1 eV. The peak at this binding energy is often reported as demonstrating evidence of bonding between carboxylic groups and the metallic surface³⁸⁻⁴⁰. The XPS Zn 2p_{3/2} spectra for EP show a peak at 1021.5 eV, corresponding to Zn/ZnO. This peak has a shoulder at higher binding energies, associated with the presence of Zn (OH)₂ and/or ZnCO₃. However, only a single component is present on the AD surface, which can be ascribed to zinc oxalate. The Zn Auger LMM peak revealed the presence of three peaks for EP sample. These peaks are related to metallic zinc and were not clear in the case of the AD surface. This might be due to the thicker zinc oxide film for the AD sample. The HR-O1s peak of samples has been fitted with three peaks: O1s(1) at a binding energy of 531.5 eV assigned to Zn-O, O1 s(2) at a binding energy of 531.6 eV assigned to Zn-OH and O1s (3) at a binding energy of 533.0 eV assigned to ZnC₂O₄/ZnCO₂. An analysis of the O1s spectra revealed an increase in Zn oxalate/carbonate for the AD sample relative to EP. The quantified concentrations associated with the C, O and Zn peaks are presented in Fig. 1h. Considering the complex nature of oxidation/precipitation on Zn surfaces, the fitting of multiple Zn species is challenging. Unreliable interpretation can be produced for the speciation of the Zn surface chemical state due to many possible Zn species with overlapping binding energies.

3.3.2 Corrosion behavior

Fig.3-2. The typical cross-sectional morphologies of EP (a, b) and AD (c, d) before and after 28 days of exposure to Hanks' solution, the surface morphologies of corroded EP (e, f) and AD (g, h) before and after removing the corrosion products and EDS analyses (i) corresponding to the points and areas highlighted on micrographs.

In order to clarify the corrosion mechanisms, the influence of the surface treatment on the polarization behavior of the AD and EP surfaces was investigated at 37°C in Hanks' solution by a morphological and chemical analysis before and after the corrosion experiment. Fig. 3-2 shows typical surface morphologies and cross sections of non-corroded and corroded samples. Cross sectional observation of corroded EP samples (Fig. 3-2 a, b) revealed the formation of a porous, non-uniform film with varied thickness from 2 to 10 μm . In contrast, the porous AD coating (Fig. 3-2c) underwent densification during the ongoing corrosion process. This may be due to blocked orifices in the anodic film from Zn degradation products, which formed a relatively dense barrier during the 28 days of exposure to Hanks' solution (Fig. 3-2d).

The effect of the surface treatment on the corrosion behavior can be more clearly assessed after removing degradation products. Signs of localized corrosion appear on the EP surface (Fig. 3-2f). Only small pits were observed on AD surface, indicating a surface attack that more than likely occurred during the deposition of the AD coating. EDS analyses of the corrosion products (which are listed in Fig. 3-2i), revealed the presence of Zn, Ca, P, O, Cl and Na. Hanks' solution often produces degradation products that contain zinc oxides, hydroxides, carbonates, calcium phosphates, and carbonates^{41, 42}. The oxygen content may imply the precipitation of oxides and insoluble hydroxides. This is a result of the ionic reaction between Zn^{2+} and several acid radicals such as Cl^- , HCO_3^- , HPO_4^{2-} , which are components of the testing solution. Corrosion products with atomic ratios of Zn:O close to 1:1 are consistent with zinc oxide, a 1:3 Zn:O ratio is typical for zinc carbonate (ZnCO_3). A ratio of Zn:O exceeding 2 might suggest the presence of zinc hydroxide. Further, the presence of ZnC_2O_4 in anodic films (for samples before and after corrosion) is confirmed by a 1:4 Zn:O ratio. Additionally, different Ca/P ratios on the degraded surfaces were found. The Ca/P ratios of various calcium phosphate salts are the following: 1 for $\text{CaHPO}_4(2\text{H}_2\text{O})$, 1.33 for octacalcium phosphate (OCP), 1.5 for tri-calcium phosphate (TCP) and 1.67 for hydroxyapatite (HA).

The electrochemical measurements provided further insights into the corrosion behavior of the surface treated samples. The respective polarization plots and their kinetic parameters are shown in Fig. 3-3a. According to the data, EP samples showed a minimum corrosion current density $I_{\text{corr}} = 17.17 \mu\text{A}$ compared to AD $I_{\text{corr}} = 20.12 \mu\text{A}$. As seen in Fig. 3a, the cathodic current densities of EP and AD samples are similar. The anodic branches showed nearly the same trend. At potentials more positive than the Tafel range, the current density plateau was established at approximately $10^{-1} \text{ A cm}^{-2}$. The anodic polarization curves of EP samples revealed a passivation-like region followed by an acceleration to a higher dissolution rate (Fig 3a).

Fig.3-3. Electrochemical measurement plots for AD and EP samples (a) potentiodynamic polarization curves (b) Nyquist and (c) Bode plots for EP samples, (d) Nyquist and (e) Bode plots for AD samples, after 1h, 24h, 72h, 120h, and 168h of immersion in Hanks' solution

The evolution of the electrochemical behavior for the EP and AD was monitored by EIS as a function of time. Nyquist and Bode diagrams for both types of zinc surfaces, EP and

AD are presented in Fig. 3-3 (b-d) for periods of exposure corresponding to 1h, 24h, 72h, 120h, and 168h. It is important to note that for 168 hours of exposure the impedances associated to the AD samples were superior to these of the EP substrate. Further, the results revealed that between 1 and 168 hours there was a large increase in the impedance of the AD samples (from $\sim 250 \Omega \cdot \text{cm}^2$ to $\sim 900 \Omega \cdot \text{cm}^2$, Fig. 3d), while that of the EP slightly decreased after 24 hours, reaching finally the value $\sim 400 \Omega \cdot \text{cm}^2$ after 168 h (Fig. 3-3b). The results of the potentiodynamic tests refers to the sample surfaces after 1h of exposure to the Hanks' solution. Accordingly, the results of EIS showed lower corrosion resistances for AD samples after 1h compared to EP (Fig. 3-3b, d). After 1h of exposure, a time constant to frequencies of around 10 kHz is observed, likely due to charge transfer processes and charging of the double layer at the Zn surface and underneath the pores in the AD film. For 24 hours of exposure, the indication of another time constant at low frequencies associated to corrosion processes at the metallic substrate is seen on the Nyquist diagram (Fig. 3-3 b). This could be explained by the localized corrosion of the EP substrate and the formation of corrosion film. When comparing Bode plots of EP (Fig. 3-3c) and AD (Fig. 3-3e), both curves in impedance and phase angle plots of anodic coating showed a shift to a lower frequency. The AD Bode diagram corresponding to 168 hours of immersion test show a shoulder at frequencies around 1 kHz and a peak at 1 Hz indicating a stable surface after the increasing dissolution of AD coating (Fig 3-3 e). Additionally, the part of AD coating under the influence of corrosive medium transformed into Zn-based oxides/carbonates/phosphates featuring low solubility and acting as a corrosion barrier. To quantify the changes in the impedance for EP and AD samples, the results presented in Fig. 3-3 were fitted using a circuit electric equivalent embedded in Nyquist plot (Fig. 3-3 b, d). The model providing an adequate fit of the EP data is shown in Fig 3-3b (inset). The EP/solution interface is interpreted as bi-layered corrosion product, made of an inner (R_i , Q_i) and an outer (R_o , Q_o) layer. The R_i and R_o of EP displayed decreasing corrosion resistance within 24 h, then increased gradually to 120 h and decreased again afterwards⁴³⁻⁴⁶. Both the capacitance and the resistance of the outer layer are high and the η value indicate a non-perfect interface (0.7–0.9) (Table S1, supplementary materials), which can be attributed to a hydrated

oxide/carbonate/phosphate layer. R_i and R_o decrease after the first 24 h, indicating the disruption of the EP surface due to the corrosion attack. After 72h, the increase in resistance is observed suggesting the growth of the degradation products. The low Q_i values after immersion times of 1 h, 24 h and 72 h are typical for porous films. Increasing Q_i after 120 h, suggest the formation of more stable, dense corrosion products layer. Accordingly, the corrosion resistance increases with degradation products formation and their accumulation on the EP sample surface. The high frequency arc on Nyquist plot was assigned to the charge transfer process, whereas the low frequency arc was allotted to the formation of unevenly distributed Zn corrosion products and species adsorbed on EP surface. The equivalent circuit depicted in Fig. 3-3d was used to fit spectra of AD sample. R_{sol} was the solution resistance, R_{ct} and Q_{dl} represented the resistance of charge transfer and capacitance of the electrical double layer; Q (constant phase element) was used here in place of a capacitor to compensate for the non-homogeneity of the system^{46, 47}. The AD film became homogeneous in terms of defects distribution, with lower number of micro-pores resulting in reduced roughness (and the actual surface area of the coating) leading to a decrease in R_{sol} . The R_{ct} value increases with progressing corrosion, suggesting the densification of the AD film. The diffusion resistance can be mainly attributed to the inner, denser layer of the coating. The reduction in Q_{ct} (reflecting the nature of the inner layer structure) comes from the thickening of the double layer, suggesting that corrosion products are formed at the surface/electrolyte interface. The rise of R_{sol} and R_{ct} indicates the formation of a compact layer on the AD surface, and charge transfer reactions in the protective film, respectively, resulting in the densification of the AD layer and the homogenized, more corrosion resistant surface structure. The solution resistance (R_{sol}) remained relatively stable during the entire experiment and the reproducibility was good for all three samples.

3.3.3 Histomorphometry and biocompatibility

A representative image of the neointimal tissue that formed around the implants is shown for each condition in Fig.3-4. Thin, non-protruding neointima is observed for the AD implants, with a variable response to the EP implants.

Fig.3-4. H&E staining depicting typical (top row) and deleterious responses (bottom row)

The following equation was used to obtain the lumen occlusion index of a sample³³,

$$LOI = \left[\frac{WLT}{BNL - \Phi} \right] \times 100\% \quad (\text{Eq. 2):}$$

The lumen occlusion index (LOI), where WLT is the wire to lumen thickness, BNL is the base neointimal length (in micrometers), and Φ is the wire diameter (in micrometers).

Fig.3-5. Quantification of in vivo histomorphometric analysis. Boxplots show the average with the boxed region ranging from 25% - 75%. Whiskers extend from 0% - 100% with each dot representing one sample point.

While there were neither failures nor reductions in biocompatibility (BC) for any of the 9 AD treated samples, the EP condition produced 1 failure based on lack of endothelialization, 2 failures based on LOI, and 2 reductions in BC, as shown in Fig.5. The reductions in BC are defined as any neointimas with a LOI $\geq 20\%$ and $\leq 30\%$ ³³. Failures are defined as any samples with LOI of $\geq 30\%$. The quantified histomorphometric analysis is shown in Fig. 3-5. One EP sample was excluded from the statistical analysis due to a lack of endothelialization. With this exclusion, no statistical difference between WLT ($45\mu\text{m} \pm 12\mu\text{m}$ vs $99\mu\text{m} \pm 100\mu\text{m}$), BNL ($813\mu\text{m} \pm 262\mu\text{m}$ vs

604 $\mu\text{m} \pm 105\mu\text{m}$) or NA ((6 $\pm$ 8) $\times 10^4$ vs (8 $\pm$ 5) $\times 10^4$) is observed. There is no difference between the two groups in terms of LOI (10 $\pm$ 5% vs 33 $\pm$ 37%), although there is a trend towards significance (p=0.118).

3.4 Discussion

Although a degradable surface film is designed to rupture and eventually dissolve, it serves as the initial interface between the medical implant and biological environment. In this central position, the surface film regulates the host response and potentially direct long-term neointimal remodeling. Although the degradation products of zinc implants have been shown to inhibit harmful neointimal responses⁴⁸, the surface film characteristics necessary to achieve optimal suppressive effects have not been explored in the scientific literature. Here, we have developed surface films through electropolishing and anodization processes with different characteristics to test the hypothesis that the surface film character of degradable zinc implants regulates neointimal responses. The character of AD and EP zinc surfaces was described based on surface features, corroded surface morphologies, cross sectional study, and electrochemical analyses. AD and EP surfaces prepared on Zn wires were then implanted into the abdominal aorta of Sprague Dawley rats for a direct comparison in terms of fundamental biocompatibility metrics.

We successfully prepared a smooth and defect-free surface (EP), similar to industry standard stents manufactured for clinical use. In marked contrast, the AD surface was rougher, with a bumpy nano-scale pattern and a thickness of about $\sim 6 \mu\text{m}$ (Fig. 3-2c). The AD surface possessed a higher atomic percentage of oxygen and carbon, with potentially a presence of zinc carbonate and oxalate. In contrast, an ultra-thin oxide film was present on the EP sample. After in vitro corrosion, a porous corrosion layer was present on the EP sample, while a densified corrosion layer was recorded on the AD sample (Fig.3-2b, 2d). After removal of the corrosion products, large areas of localized corrosion were observed on the EP surface, while only small defects were visible on the AD surface that are ascribed to the surface attack during anodization process (Fig. 3-2f,

2h). Although the EP and AD surfaces possessed similar corrosion potential values with similar corrosion rates (Fig.3-3a), the anodic region of the EP curve on Fig.3-3a displays a tendency towards dissolution when compared to AD, revealing a slight passivation of the AD surface. The AD surface was resistant to corrosion overall, as shown by the initially high impedance values on the comparable Nyquist plots (Fig.3-3b, 3d) and the densified corrosion layer that formed on the AD sample. In contrast, the Nyquist and Bode plots of the EP condition revealed inferior corrosion resistance. Indeed, the development of localized corrosion was confirmed from the 28-day immersion in Hanks' solution (Fig.3-2f).

Fig.3-6. Schematic illustration of corrosion process of AD and EP samples.

The contributions of the prepared surface films to the corrosion behavior of the Zn materials can be described based on the presented in vitro pre and post corrosion analysis. With increasing exposure time, the reaction mechanism for carbonates, phosphates, and hydroxides is significantly slower, and mutual diffusion from the Zn substrate and Hanks' solution is reduced. Principal diffusion pathways are provided by the capillaries in the porous corrosion layers (and anodic film in the case of AD samples, Fig.3-6), where Zn ions diffuse from the bulk material toward the solution/surface interface. Ions present in the body fluid diffuse in the opposite direction (shown in Fig. 3-6). Thus, the degradation rates are reduced with corrosion product thickening and its densification.

The surface oxide-hydroxide-phosphate-carbonate film on AD and EP samples cycle through a process of partial dissolution and re-precipitation in Hanks' solution. When the dissolution rate is larger than that of re-precipitation, ions are gradually released. When the metallic surfaces are covered with a thick and compact film, the corrosion resistance increases. However, the composition of this protective film might change even though these precipitates are macroscopically stable. The AD samples exhibited a higher corrosion resistance compared to EP, during longer exposure to Hanks' solution as shown by EIS measurements (Fig. 3-3). The Zn degradation precipitates blocked pores and capillaries in the AD film, protecting the Zn metal during longer exposures to physiological solution. In contrast, the protective film formed on EP samples was less compact, with varied thickness. Thus, it provided limited corrosion protection.

Since the noted differences in surface films influence corrosion behavior, we asked whether the different oxide film characteristics could provoke changes in the biocompatibility of zinc implants in the abdominal aorta of rats. Based on nine observations per treatment group, the AD surface produced stable, non-obstructing neointimal growth, while the EP preparations evoked a high rate of harmful responses (Fig. 3-4). Specifically, we observed two reductions in biocompatibility for the EP condition, and two outright failures in terms of $\text{LOI} \geq 30\%$ (Fig.3-5). Another failure of one EP specimen was due to lack of endothelialization. The lack of roughness in the EP preparation vs. AD may explain the delay in endothelialization, as 3-dimensional surface features may aid in cell adhesion and migration^{49, 50}. In total, three out of the nine observations failed for the EP condition, and an additional two had a reduction in biocompatibility. In contrast, the nine AD specimens were free from failures or reductions in biocompatibility.

When evaluating the biocompatibility of vascular materials that are destined for human use, the numerical clustering of the BC metrics is an important consideration, even when the average response is acceptable from an application standpoint. The large spread in the tabulated data for the wire to lumen thickness (WLT) and lumen occlusion index (LOI)

for EP specimens raises serious concerns, as it suggests that the in vivo response is non-uniform. The variability in biocompatibility responses could be a consequence of the non-uniform in- vitro corrosion behavior. Critically, corrosion product stability is reduced on the EP surface, leading to localized corrosion and pitting. The increase in substrate attack (apparent in Fig. 3-2) caused by a porous, loosely compacted corrosion layer that aids in solute diffusion to the metal-product interface could promote localized sites of inflammation along the wire that initiate neointimal growth and smooth muscle cell proliferation. Thus, increased variability from localized corrosion due to surface film character directly translates to impaired in vivo performance. Overall, the EP condition increased variability in the WLT metric, and produced a near significant increase in the LOI ($p=0.118$). The EP surface produced a 22% failure rate based on the LOI exclusion criteria (2 out of 9), with only 44% of the observations meeting the full biocompatibility benchmarks (4 out of 9). 100% of the AD samples (9 out of 9) met the full biocompatibility benchmarks, leading to a failure rate of 0 % (0 out of 9). To provide context with clinical observations of stent performance, bare metal stents incur a restenosis (failure) rate of between 20%-30%, with drug eluting stents reducing this value to <10%. Using the restenosis rate of present generation drug eluting stents (~10%) as a guideline, the AD surface is seen as potentially improving outcomes while the EP surface falls outside expectations for success. Although the AD coating employed here may not be appropriate for stents due to potential cracking of the layer during balloon crimping and deployment, this investigation highlights the critical importance of oxide films for determining the biocompatibility of degradable zinc metal implants.

3.5 Conclusions

An electropolished and anodized surface oxide film with widely varying characteristics were prepared on degradable zinc implants and compared, both in vitro and in vivo, to clarify the relationship between surface film properties and biocompatibility. The following conclusions were reached:

The anodized samples exhibited a higher corrosion resistance compared to electropolished during longer exposure to the physiological solution due to the densification of the oxalate film, which acted as a barrier limiting the contact of the substrate with electrolyte.

The corrosion product film formed on electropolished samples was less compact, with varied thickness. Thus, it provided limited corrosion protection.

The electropolishing produced a smooth and thin oxide film surface that elicited a high variability in biocompatibility, with a high rate of failure, possibly due to surface corrosion heterogeneity.

In contrast, the corrosion stability of the anodized surface film corresponded to improved biocompatibility outcomes.

In particular, the ability to tune the in vivo biodegradation behavior and biocompatibility of the Zn biomaterial as a function of surface properties was experimentally verified

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3.8 References

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4 In-Vivo Biocompatibility of Next Generation Zn-Ag Based Stent Materials ⁷

4.1 Introduction

Since the introduction of zinc as a degradable metallic candidate for bioresorbable stents in 2013[1], numerous zinc alloy formulations with improved mechanical properties and microstructure relative to pure zinc have been investigated [2-5]. A central feature of the zinc-based stent materials being developed is their gradual degradation in the arterial environment, until their eventual disappearance [6, 7]. The biodegradable feature of zinc-based materials makes them highly interactive with the host environment. Although bioresorbable implants differ considerably from conventional biostable implants in terms of their interaction with the host, most studies provide limited details regarding the biological effects evoked by the implant's presence and degradation. Consequently, there remains a wide deficit in our understanding of the impact of alloying elements and bulk processing on the tissue response to zinc-based implants. The biointeractivity of zinc-based materials needs to be thoroughly characterized in order to engineer zinc-based implants with improved safety and efficacy relative to their bioinert metallic counterparts.

Our group has worked to clarify the relationship between the material composition, properties and surface finish of degrading zinc implants and the biologic responses they evoke in the arterial environment [8-15]. We make use of an implant model wherein a wire made of the material in question is implanted into the abdominal aorta of healthy Sprague Dawley rats and allowed to degrade for 3-20 months[9, 12, 13]. Recently, we developed a histomorphometric approach to quantify the performance of bioresorbable materials in the arterial environment [11]. This work has identified thresholds that distinguish between acceptable materials and those that stimulate a harmful smooth muscle cell proliferative response, a hallmark failure of long-term implanted stents.

⁷ Material in this chapter is in preparation to be submitted to a journal for publication

Of all the zinc-based implants evaluated so far, pure zinc generally evokes the most favorable response within the vascular environment. A stable neointima forms around pure zinc that is restricted in terms of area, thickness, and progression and generally avoids inflammatory infiltrates for up to 11 months. We recently reported that alloying with magnesium contributes to the biological failure of zinc implants through excessive tissue growth [11]. Although the implantation of a Zn-4Li alloys into the arterial environment did not produce outright failures, this alloy experienced a reduced biocompatibility metric relative to pure zinc due to an increased degree of inflammation. In addition to modifications to the bulk matrix, we have also shown that the initial surface oxide film characteristics dramatically affect the early neointimal growth [9][16]. For instance, thin surface oxide films produced by conventional electropolishing are susceptible to large variations in corrosive attack that lead to excessive variations in biological responses[9]. In contrast, relative stability of the surface oxide film (imparted for example through anodization) promotes a more homogenous corrosion along the length of the wire, which results in more positive and reproducible biological responses. In contrast to some success seen in modifications of the surface[9, 16], manipulating the bulk metal through alloying to improve the mechanical properties has tended to reduce the overall biocompatibility of the implant material relative to pure zinc.

As metallurgical modifications continue to advance and alloy compositions become more complex, it has become more urgent to clarify the relationship between material characteristics and tissue response. Such clarifications may provide predictive power for metallurgists to aid in the design of zinc-based medical implants. More recent zinc systems include 2-4 alloying elements to refine the grain structure, increase microstructural stability, and improve the mechanical properties of zinc to acceptable levels for stenting applications [2, 4]. The addition of each element impacts corrosion behavior and products, secondary phases within the microstructure, and surface characteristic, all of which can affect the overall device biocompatibility at various stages after implantation. A deeper understanding of the relationships between material modifications and biological responses is needed to identify desirable material

characteristics from a biocompatibility standpoint, providing a foundation for metallurgical engineering of bioresorbable medical alloys.

Recently, a new Zn alloy system has been introduced with the addition of Ag[17]. This system benefits from an increased volume fraction of AgZn_3 particles that precipitate at the grain boundary and provide grain boundary strengthening due to impaired propagation of dislocation slip planes, as well as increased grain refinement (grain size $1.5\text{ }\mu\text{m}$)[18]. The Ag addition dramatically increases the ultimate tensile strength to 280-290 MPa without sacrificing elongation to failure, and provides a beneficial antibacterial effect [2, 18].

Here we leverage high sample sizes to investigate the biocompatibility of 3 novel Zn based Ag-containing alloys, ranging from binary to quinary alloy systems. This work was intended to clarify the relationship between elemental profile and biocompatibility for the Zn-Ag system. Selected binary and quinary Zn-Ag-based alloys underwent thermal treatment (TT) to increase the solubility of Ag-rich phases within the Zn bulk matrix, yielding 2 different microstructures with the same elemental composition. A side-by-side comparison of implant materials with and without TT was intended to clarify the effect of microstructural features on biocompatibility. The insight gained from these experiments is needed in order to integrate biological considerations into the metallurgical design of fine/ultrafine-grained structure Zn-based materials through alloying and processing.

4.2 Methods

4.2.1 Alloy Formulation

High-purity starting metals (>99.99%) were melted at 700°C in a sealed cylindrical graphite crucible in an electric resistance furnace to minimize oxidation and ensure that sound castings are produced. The ingots had a cylindrical shape with a nominal diameter of 28 mm and a length of 100 mm. The bars were machined to a nominal diameter of 25 mm and homogenized for 8 hours at approximately 400°C followed by water quenching.

Nominal elemental formulations were used as outlined below:

Zn-4Ag (wt%) (binary)

Zn-4Ag-0.6Mn (wt%) (ternary)

Zn-4Ag-0.8Cu-0.6Mn-0.15Zr (wt%) (quinary)

The homogenized alloys were subsequently extruded at 310°C with an extrusion rate and speed of 39:1 and 0.5 mm/min, respectively, to obtain cylindrical rods with a diameter of 4 mm. The extruded rods were cold drawn at Fort Wayne Metals (Fort Wayne, IN) to a wire with a diameter of 0.25 mm.

The binary and quinary alloys were heat treated in order to minimize the Ag-rich precipitates. For the binary alloy, TT was performed at 360°C for 10s, while the quinary alloy, due to its multiplicity of alloying elements, required higher temperature and soaking time (390°C for 20 min).

4.2.2 TEM preparation

Transmission electron microscopy (TEM) samples were prepared using standard routes, including grinding and polishing the wires to < 40 mm in thickness followed by ion beam milling to electron transparency. Low angle-ion beam milling was performed at liquid nitrogen temperature by a GATAN precision ion polishing (PIPS) II using Ar⁺ initially at 5 keV and final milling at 1 keV. TEM data were obtained with JEOL JEM F200 transmission electron microscopy operating at 200 kV.

4.2.3 Implantation and collection

Implantations, implant retrieval, and histological processing were performed as previously described [11]. Wires were implanted into female Sprague Dawley rats for 6 months. Sixteen samples were produced for each material.

4.2.4 Statistical analysis

All data was tabulated in Excel 2016 and uploaded to MATLAB 2016Ra. Raw data was transformed into log space for statistical comparisons. An unpaired two sample Welch's t test was used to investigate the experimental groups. When appropriate, an F test was used to identify differences in the variance. All data was approximately normal.

4.3 Results and Discussion:

4.3.1 Microstructural Characterization

Fig. 4-1. (a, b) TEM images of as-drawn binary and quinary alloys, (c, d) their corresponding EDS elemental

Fig. 4-1 shows the transmission electron microscopy (TEM) images of the binary and quinary alloys before and after the thermal treatments. It can be clearly seen from Fig. 4-1a and b that for both alloys the wire drawing results in the formation of an ultrafine-grained (UFG) structure (grain size of 600 nm and 200nm for binary and ternary alloys, respectively). Formulated alloys underwent tensile testing and microstructure analysis. However, because mechanical properties of alloys are irrelevant to biocompatibility study they will be presented in a separate publication. Only selected mechanical data are provided in this section to guide general correlations between microstructure and mechanical properties of the alloys.

Fig. 4-1 shows the transmission electron microscopy (TEM) images of the binary and quinary alloys before and after the thermal treatments. For both alloys the wire drawing resulted in the formation of an ultrafine-grained (UFG) structure with grain size of 600 nm and 200nm for binary and ternary alloys, respectively (Fig. 4-1a&b). An EDS elemental map of the as-drawn binary alloy wire demonstrates the formation of submicron deformation induced-AgZn₃ precipitates within a size range of 100 nm - 500 nm (red particles) uniformly dispersed in the Zn matrix (Fig. 4-1c). The formation of these precipitates markedly enhances the ductility such that the material exhibits exceptional room temperature superplasticity (fracture elongation > 400%). However, this superplasticity is accompanied by a remarkable loss of mechanical strength (~ 73%). In the case of the quinary alloy (Fig. 4-1b and d) the considerably smaller grain size relative to the binary alloy (200 nm vs 600 nm) is attributed to the extremely refined and uniformly dispersed MnZr-rich particles. Such particles hinder the growth of the newly recrystallized UFG grains during the drawing process, resulting in markedly smaller grain size. The EDS elemental map of the quinary alloy shows that the addition of Cu appreciably decreases the fraction of AgZn₃ precipitates (Fig. 4-1d). This results in a lower fracture elongation of the quinary relative to the binary alloy (90% vs 430%). To restore the mechanical properties, thermal treatments were carried out on the as-drawn alloys. Fig. 4-1e shows that for the binary alloy, TT at 360°C for 10s leads to dissolution of a large number of the AgZn₃ precipitates as well as a significant grain growth. In fact, the remarkable decrease in the

volume fraction of AgZn_3 precipitates in TT binary alloy could restore almost 73% of the mechanical strength. In the case of the quinary alloy, TT was conducted at a significantly higher temperature and for a longer soaking time (390°C , which corresponds to 96% of the alloy's melting point, for 20 min). Fig. 4-1f reveals that, surprisingly, despite the very high temperature, only a small grain growth occurs (from 200 nm to $2.4\text{ }\mu\text{m}$). This is attributed to the greatly fine and uniformly distributed MnZr-rich intermetallic particles at grain boundaries, which are outstandingly thermally stable, that prevent grain coarsening even at high temperatures. Indeed, the TT quinary alloy could substantially restore its mechanical strength (around 84%) while exhibiting a high fracture elongation of 32%.

4.3.2 General Histologic Presentation

Fig.4-2. Representative H&E and VVG stains of the binary, ternary, and quinary alloys. The neointima is outlined in yellow to increase clarity. Scale bar is approximately $500\text{ }\mu\text{m}$

The general histological presentation of the tested alloys is shown in Fig. 4-2. Typical binary Zn-4Ag, ternary Zn-4Ag-0.6Mn, and quinary Zn-4Ag-0.8Cu-0.6Mn-0.15Zr implants are shown with H&E and VVG stains. H&E staining is ideal for microscopic cell identification and characterization of the general tissue reaction, while the VVG technique clearly differentiates elastic lamina fibers and neointimal tissue. Since we are

mainly concerned with the macro neointimal development around each implanted alloy, the VVG stain is primarily used for ease of histomorphometric measurements.

The binary and ternary alloys display excessive neointmal thickness and area, with the quinary alloy showing a reduced, ideal neointimal type at 6 months. The binary and ternary alloys show evidence of increased inflammation within the neointima, by H&E identification of inflammatory cells (not shown). This was largely absent for the quinary alloy, with a reduced cell density and stable neointima growth that does not appear to be progressive.

4.3.3 Effects of alloying

Fig.4-3. Boxplots of Pure Zn, copper-free alloys, and Cu containing alloys for histomorphometrics. Panels A and B show comparisons using a Welch's two sample t test. Panels C and D show comparisons of the variance with a two sample F test.

The effects of alloying and processing on the in vivo performance of degradable zinc materials was compared in terms of histomorphometry parameters, using an approach previously described [11]. Fig. 4-3 shows a high sample size pooled performance of alloys implanted from 3-12 months, including pure zinc, multiple copper-free alloy

systems (Zn-4Ag, Zn-4Ag-0.6Mn, Zn-4Li, and Zn-XMg [0.1,0.001,0.001wt%]), and the Zn-4Ag-0.8Cu-0.6Mn-0.15Zr alloy developed by our group. In vivo performance was quantified by neointimal thickness (WLT) and neointimal area (NA). Interestingly, a similar WLT was found between the copper-free alloys and the quinary system relative to pure zinc (Fig. 4-3A). However, there is a clear superior performance of the copper-containing quinary alloys relative to the copper-free alloys, ($P=0.000019$).

The copper-free alloys generally incur a reduced biocompatibility relative to pure Zn in terms of NA, with a higher sample mean ($P=0.0016$). The NA for the copper-containing alloy is significantly reduced relative to the copper-free alloys ($P=0.0000000035$), and even improved over that of pure zinc. We attribute the reduction in NA to a reduced inflammation at the interface of the Cu containing alloy (Fig. 4-2).

A previously described lumen occlusion index (LOI) was used to compare novel alloys in terms of negative proliferative responses that occludes the lumen. Surprisingly, none of the new alloys exceeded the biocompatibility limit LOI value of $\geq 30\%$, or the reduction in compatibility limit LOI of or $\geq 20\%$. The excellent LOI values for the Zn-Ag alloys may be due to their extended base neointimal length (BNL), along with typical WLT values (which could indicate positive remodeling). Although none of the implants failed outright, the large NA produced by the binary and ternary alloys is concerning. This large increase in area is primarily due to an increase in inflammation, which is a well-known contributor to negative stent outcomes. The LOI measurement does not take into account NA, but rather relates to the protrusive growth of the neointima into the luminal space. Our lab has found that a neointimal protrusion metric is generally related to smooth muscle cell proliferation, while a large increase in NA is generally indicative of excessive inflammation[11]. Both factors should be considered when gauging the biocompatibility of degradable materials, especially since satisfactory performance on the LOI scale does not demonstrate optimal performance in all histological metrics.

Sample variance between the experimental materials is a major consideration when evaluating histological samples. Tighter data spread is associated with more uniform

performance. This feature is especially valuable in the intra luminal environment, which is subject to occlusion at any point along the length of the implant [9]. Pure zinc generally exhibits an ideal histological variance, avoiding corrosion behavior that provokes large changes in neointimal growth along the length of the implant. While there are generally no differences in the variance of WLT between pure zinc implants and the copper-free alloys, the copper-containing quinary alloys demonstrate a much tighter data spread than copper-free alloys ($P=0.029$, Fig.4-3c). NA measurements for the copper-free alloys are generally more variable relative to both pure Zn ($P=0.0044$) and copper-containing alloys ($P=0.0238$). The quinary alloy and pure Zn exhibit a similar variance, suggesting a more ideal performance from copper-containing alloys in terms of generating consistent neointimal morphometry.

Elemental composition is a serious consideration when designing alloys. When the binary Zn-Ag composition was alloyed with Mn, the biocompatibility (BC) was considerably reduced. The BC then recovered following the addition of Zr and Cu. While it can be difficult to isolate the effects of the Mn addition (which can alter corrosion behavior and form new inter metalics), it is apparent from our results that changes in the elemental profile of a degradable zinc implant can dramatically impact the BC.

4.3.4 Effects of nano-precipitate phases

Fig.4-4 Three representative VVG stained sections of the binary and quinary alloys. High variability in neointimal size can be seen in both the binary and its solution treated counterpart. A decrease in overall NA is seen in the quinary alloy, with an even greater reduction in its solution treated counterpart. Scale bar is approximately 500 μm .

In order to clarify the effects of nano-precipitate phases on neointimal growth, the binary and quinary Zn-Ag alloys were thermally processed to dissolve the insoluble deformation induced-AgZn₃ nano-precipitates. Applying TT has the benefit of reducing the amount of precipitates and thereby restoring the mechanical strength of the alloy [19].

Fig.4-4 shows the in vivo performance of both as drawn and solution treated binary and quinary alloys. For each alloy, three representative samples are depicted to demonstrate typical responses of the alloy system. For the binary system, large sample-to-sample differences are present in the NA (outlined in each image). The variability between samples does not appear to change after solution treatment (Fig.4-4 second row). The quinary system shows a marked reduction in NA relative to the binary system, along with a decreased variance. Solution treatment of the quinary alloy further reduces the NA and WLT, demonstrating the benefit of the solution treatment to biocompatibility.

The complete histomorphometric analysis of these samples is shown in Fig. 4-5. As shown in Figs 4-5A and 4B, WLT (but not NA) slightly benefitted from the solution treatment of the Zn-Ag binary alloy. The Zn-Ag and Zn-Ag TT materials both evoked an increased inflammation relative to pure zinc, which we attribute to the presence of Ag precipitates. It is well known that Ag nanoparticles contribute to inflammatory phenotypes in macrophages and elicit direct cell toxicity [20, 21]. Therefore, a more pronounced reduction in Ag precipitates (obtainable by increased treatment time or temperature) may be beneficial in terms of biocompatibility.

The quinary alloy containing Ag and Cu possesses the same Ag nano precipitates. Interestingly, after thermal treatment of the Zn-Cu containing alloy, in vivo performance was dramatically improved, and even superior to pure zinc. WLT ($P=0.023$) and NA ($P=0.000028$) from the Zn-Cu-TT alloy readily outperforms pure zinc. It is interesting that the solution treatment for the binary alloy did not improve the BC as dramatically as for the quinary system. The solution treated binary alloy microstructure retains

substantial Ag precipitates, possibly a consequence of the low ST time that was chosen in order to optimize the mechanical properties. Although the untreated quinary alloy also possesses Ag nano-precipitates, we speculate that the beneficial cellular effects from the addition of Cu overwhelms the negative response that is generated from these harmful phases.

Fig.4-5. Comparisons of the histomorphometric performance of binary Zn-Ag

and quinary Zn-Ag-Mn-Zr-Cu alloys in the as-drawn form and after solution heat treatment. All comparisons were made with a two sample Welch's t test.

We are the first to report a zinc-based alloy (Zn-4Ag-0.8Cu-0.6Mn-0.15Zr (wt%)) that performs better than pure zinc in terms of biocompatibility. We speculate that the Cu inclusion has a beneficial impact on inflammatory development and neointimal growth. Cu has been shown to increase the proliferation of endothelial cells, but not smooth muscle cells [22]. In a proinflammatory environment such as a newly injured artery, increased scavenging of reactive oxygen species could have a beneficial effect on neointimal progression and inflammation. Cu can also directly act as a catalyst for the generation of NO from endogenous RSNO's, which could also inhibit inflammation [23] and SMC growth [24]. Furthermore, increased superoxide dismutase (SOD) activity in EC's could promote a favorable balance towards additional NO release [25].

4.4 Conclusions

The findings of the study conclude:

- Addition of alloying elements to zinc usually decreases the biocompatibility of zinc implant in vascular environment by increasing inflammation responses;
- Ag-rich nano-precipitates decrease biocompatibility, a phenomenon that can be counteracted by dissolving the AgZn₃ precipitates in the bulk Zn matrix;
- Cu addition to Zn-Ag alloy reduces inflammation and substantially increases biocompatibility, even in the presence of harmful AgZn₃ precipitates, an effect that is enhanced when the precipitates are dissolved in the Zn matrix; and
- Neointimal forming cells are responsive to elemental additions and microstructural changes in degradable zinc based materials.

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5 Zinc Based Materials Regulate the Vascular Biologic Response to Implants by Releasing Therapeutic Transition Metal Ions⁸

5.1 Introduction

Zinc (Zn) based degradable scaffolds under development for cardiovascular applications have garnered increasing attention in recent years [1-5]. Metallurgical innovations that increase the mechanical properties have positioned this class of degradable metals to move forward in preclinical testing of stent prototypes [6]. However, while there has been extensive research and development with regards to alloying and materials refinement, the biological consequences of a degrading zinc-based arterial implant have not been well addressed in both fundamental and clinical studies.

The concept of biodegradation is straightforward; physiological fluids break the material down into byproducts that are either metabolized or cleared from the implant site. As the implant degrades, the space occupied by the material is gradually infiltrated by cells and matrix and reintegrated into the host environment. As with all implanted materials, the initial cells arriving at the material-tissue interface are predominantly macrophages and foreign body giant cells that help clear the implanted material [7]. For arterial implants, smooth muscle cells (SMCs) eventually form a dominant cell type within a developing neointima [8]. For materials that are designed to degrade, an added consideration beyond the original material is the effect of degradation byproducts on the behavior of infiltrating cells. Magnesium (Mg) alloys and polymers such as polylactic acid (PLLA) remain the most widely studied materials for fully degradable stent devices in terms of the biological action of byproducts. Magnesium degrades into Mg^{2+} , MgO, and $Mg(OH)_2$, eventually leaving behind a Ca/P rich product [9-11]. PLLA degrades by hydrolysis into nontoxic biocompatible products [12]. Similar to the selection of bioinert metals for conventional

⁸ Material in this chapter is in preparation to be submitted to a journal for publication

biostable stents, a primary consideration behind selecting bioresorbable materials for stenting has been to avoid negative biological responses. Indeed, Mg and PLLA materials have advanced through pre-clinical studies and entered into clinical trials in large part because their degradation byproducts do not interfere with the normal functioning of the native surrounding cells and are highly unlikely to promote systemic effects[11-13].

Because the byproducts of degradation can diffuse away from the implant interface, clarifying the biological action of degradation byproducts on vascular smooth muscle cells is of paramount importance. Early work with zinc uncovered LD₅₀ values that were well below that of Mg and Fe (iron) [14, 15]. This raised the prospect that ionic Zn might exert a strongly toxic effect on arterial cells. However, the exact opposite way of thinking was suggested by early in vivo findings. Low inflammation, the absence of any overt signs of necrosis, and a stable neointima predominated around Zn implanted into rat arteries out to ~6 months[16]. Based on the in vitro findings, one would have expected to see a strong inflammatory response, the clear presence of necrotic regions, and a negative neointimal progression. Together, the in vitro and in vivo findings suggest that the biological effects of zinc on neointimal progression may not be mediated by toxicity, but rather by programmed intracellular signaling pathways.

Due to zinc's inherently low mechanical properties, a Zn stent material requires the addition of alloying elements[1-5]. Alloying of zinc with non-toxic elements at their specified concentrations can be used to improve mechanical properties and refine microstructure of the implant. If an element exerts beneficial bioactive effects on arterial cells, its incorporation into the bulk matrix ensures its elution as the implant degradation progresses. In such a manner, an advanced bioresorbable stent metal can be engineered to release multiple beneficial bioactive elements at optimized concentrations for the lifetime of the implant. Alloying with copper, for instance, can improve the structural and mechanical properties of zinc implants[17]. In this communication, we introduce a new concept of using Zn²⁺ and Cu²⁺ ions as potential therapeutic elutants that are supplied to a host site from degradable zinc-based implant.

5.2 Experimental

5.2.1 Materials

ACS reagent grade methanol was purchased from Millipore Sigma (St. Louis, MO). Bradford assay kit (ab102535), caspase-3, caspase-8 and caspase-9 multiplex activity assay kit fluorometric (ab219915), TUNEL assay kit (ab206386), anti-cleaved caspase-3 antibody (ab2302), anti-alpha smooth muscle actin antibody (ab5694), and goat anti-rabbit IgG Alexa Fluor 488 were all purchased from Abcam (Cambridge, MA). 4',6-Diamidino-2-phenylindole dihydrochloride (DAPI) (D8417), Dulbecco's modified eagles medium-low glucose (DMEM) (D6046), Dulbecco's phosphate buffered saline (PBS) (D8662), penicillin-streptomycin (P4333), goat serum (G9023), zinc acetate (383317), D-glucose (G8270), and fetal bovine serum (F2442) were purchased from Sigma-Aldrich (St. Louis, MO). 3-morpholin-4-ylpropane-1-sulfonic acid (MOPS) (172630250) was purchased from Fisher Scientific (Toronto, ON). High purity Zn wires (4N-99.99wt%) were purchased from Goodfellow (Coraopolis, PA). Platinum wires were purchased from Fort Wayne Metals (IN).

5.2.2 Methods

5.2.2.1 *In vivo Implantation*

Pure zinc (Zn) and platinum (Pt) wires were implanted into the abdominal aorta of adult female Sprague Dawley rats for 3 months, n=5. A detailed description of the surgical procedure was provided in our earlier publication [18]. The wire and tissue explants were collected, surrounded by optimal cutting media, and snap frozen in liquid N₂. The samples were stored at -80°C until cryo-sectioning.

5.2.2.2 *General Histology, immunofluorescence and in-situ apoptosis detection*

Wire/tissue samples were cross-sectioned with a cryo-microtome at 10 μ m section thickness and placed on histobond glass slides. Multiple sections spanning a ~0.2 mm distance were collected for each sample. For hematoxylin and eosin staining (H&E) and TUNEL labeling, sections were fixed in paraformaldehyde for 10 min. H&E staining was carried out by protocols described by the authors previously [18, 19], and TUNEL labeling was performed following manufactures' protocols.

For immunofluorescence, sections were fixed in ice cold methanol for 5 min and allowed to air dry before washing. The slides were rinsed 3X in PBS, then blocked in a 10% (v/v) goat serum solution in PBS for 30 min. After blocking, slides were either incubated with an anti-cleaved caspase-3 antibody (1/100 dilution) overnight at 4 °C, or an anti- smooth muscle alpha actin antibody (1/250 dilution) for 1 hr at room temperature (RT). After incubation, the slides were rinsed 3x with PBS and incubated with goat anti-rabbit IgG Alexa fluor 488 (1/300 dilution) for 1 hr at RT. The secondary solution was rinsed with PBS 3x, then stained with a DAPI solution (1/1000 dilution of 1mg/mL stock) for 2 min at RT. DAPI was rinsed off with PBS and the slides were mounted with a glass coverslip and aqueous mounting media, and imaged immediately with an upright BX51 Olympus epifluorescence microscope.

5.2.2.3 *Ex-vivo arterial ring tissue culture*

In order to investigate the potential of Zn^{2+} to activate caspase-3 in smooth muscle cells directly, an ex vivo arterial culture model was developed. The thoracic aorta was collected from donor adult female Sprague Dawley rats. Each thoracic aorta was cut into rings of approximately 5-8 mm in length in Dulbecco's PBS supplemented with 1% penicillin-streptomycin (PS) (v/v). The arterial rings were randomized and allowed to recover overnight in low glucose DMEM supplemented with 10% FBS and 1% PS, at 5% CO_2 and 37°C. After overnight recovery (to allow cells to adjust to culture conditions and recover from injury), the arterial rings were randomly immersed in a physiological saline solution (131 mM NaCl, 4 mM KCl, 2.5 mM $CaCl_2$, 5.5 mM glucose, and 1 mM $MgCl_2$)

buffered by 10 mM MOPS at a pH of 7.25 (pH adjusted with sodium hydroxide), or the same physiological saline solution with the addition of 0.5 mM Zn acetate, MgCl₂, FeCl₃, or CuCl₃. The arterial rings were incubated in the control and insult loaded saline solutions for 12 hours.

After incubation, the smooth muscle cell containing media compartment was mechanically separated from the other arterial compartments. The adventitia compartment for each arterial ring was bluntly dissected from the media layer using forceps under a stereomicroscope. Removal of the endothelial cells was confirmed with en-face DAPI imaging (not shown), and the media layer was immediately washed in PBS containing no magnesium and calcium and stored in lysis buffer at -80°C for further processing.

5.2.2.4 *Multi-caspase activity assay*

Caspase-3, caspase-8, and caspase-9 activity was detected for the media compartment of the arterial rings using a plate based fluorescent assay. The compartment was homogenized in a lysis buffer provided by the manufacturer, and protein determination was carried out using a Bradford assay kit. The multiplex caspase activity kit (abcam ab219915) provides each substrate used to detect its respective caspase in a conjugated form to fluorophores with wide spectral separation, allowing multi-caspase activity to be determined in the same lysate. There was a minimum of 4 samples per condition (n=4).

5.2.2.5 *Confocal microscopy*

Confocal microscopy was used to inspect the *in situ* localization of active caspase-3 within the zinc-insulted arterial segments. After collection, thoracic aortas were allowed to recover in supplemented DMEM overnight at 37°C. The segments were then rinsed in PBS and incubated in either a 0.5 mM zinc solution or control saline solution. The segments were then rinsed in PBS, surrounded by optimal cutting medium and snap frozen in liquid N₂. 50 µm thick cryosections were taken at five evenly spaced locations. These sections were stained for active caspase-3 as described above, imaging three

locations per section with an Olympus FluoViewTM FV1000 laser scanning confocal microscope. 30 μm volume stacks were taken for each 600x magnified field.

5.2.2.6 Corrosion of alloys and ICP-OES analysis

Ternary Zn-4Ag-0.6Mn and quinary Zn-4Ag-0.8Cu-0.6Mn-0.15Zr ultrafine zinc wires were corroded in Hanks balanced salt solution [20] for 14 and 28 days. The solutions supernatant were then sterile filtered with a 0.22 μm syringe filter. After filtration, the solutions were acid digested with trace metal basis nitric acid, and analyzed using inductively coupled optical emission spectroscopy (ICP-OES). The detection limit for copper is 10 ppb.

5.2.2.7 Statistical Methods

An unpaired two sample student's *t* test with unequal variance was used to evaluate experimental groups. Error bars are given as sample standard error. A minimum of 4 sample replicates ($n=4$) was included for all experimental groups. A one-way ANOVA was used to compare multiple groups.

5.3 Results

5.3.1 Morphometric presentation

Fig.5-1 General histomorphometry and neointimal cellularization of Pt and Zn

The typical appearance of the neointimas (NI) that developed on Zn and Pt wires is shown in Fig.5-1. H&E staining at 100x normal magnification depicts thin, continuous NI formation over both wires. In order to compare the cellularity of the NI produced by the two materials, DAPI staining was performed, as shown in the bottom two panels of Fig.1. Both NIs appear to be cellularized with a similar spatial distribution of cells. Furthermore, histomorphometric parameters, including NI area for Pt vs. Zn, $(5.4 \pm 1.6) \times 10^4 \mu\text{m}^2$ vs. $(6.6 \pm 1.9) \times 10^4 \mu\text{m}^2$ and

wire to lumen thickness ($50 \pm 12 \mu\text{m}$ vs. $59 \pm 21 \mu\text{m}$) show a slight but non-significant elevation for Zn ($p=0.308$ and $p=0.433$, for area and thickness, respectively) ($n=5$ per condition).

5.3.2 TUNEL labeling of NI and α -SM / active caspase-3 presence

Cross sections of the Pt and Zn implants were labeled for TUNEL+ identification. The Pt specimens displayed low TUNEL+ staining in both the luminal and mural regions of the NI. In stark contrast, all Zn specimens demonstrated strong TUNEL+ staining within both the mural and luminal portions of the NI. When taken as a percentage of area covered by threshold analysis, the NI surrounding Zn implants was significantly more TUNEL+ ($2.4 \pm 2.1\%$ vs $6.5 \pm 1.5\%$, $p=0.008$, Pt $n=5$ vs Zn $n=5$, respectively).

In addition to higher TUNEL+ staining within the NI surrounding Zn implants, we observed a reduced amount of α -SMA area coverage in Zn NI tissue compared to the platinum control ($23 \pm 2\%$ vs $14 \pm 6\%$, $p=0.017$, Pt vs Zn, respectively). This reduced amount of α -SMA area coverage for Zn specimens was associated with an increased amount of area coverage in the NI for active caspase-3 ($4.3 \pm 2.7\%$ vs. $8.6 \pm 2.0\%$, $p=0.032$, Pt $n=5$ vs Zn $n=4$ respectively).

Fig.5-2 Neointima cellular and protein expression of Pt and Zn. TUNEL, alpha smooth muscle actin, active caspase 3 and DAPI staining were performed for a minimum 3 sections per sample (n=5 samples per condition). Data error is reported in standard sample error and statistical significance is shown by a Welch's T test. Scale bar is set at approximately 100µm.

5.3.3 Dose dependent caspase -3, -8, and -9 activation by Zn

In order to determine whether Zn may promote caspase activation in smooth muscle cells, we incubated arterial segments in multiple concentrations of Zn for 12 hours, ex vivo. We found that smooth muscle cell caspase-9 is not activated by 0.5 mM Zn (Fig 5-3a). Interestingly, higher Zn concentrations significantly suppress caspase-9 activity from basal levels (1 mM and 1.5 mM, $p < 0.00005$). An increase in caspase-8 activity is detected at 0.5 mM relative to the control condition, Fig.5-3b ($p < 0.0005$). Higher concentrations of zinc significantly inhibit caspase-8 relative to the 0.5 mM condition (1 mM and 1.5 mM, $p < 0.00005$). Caspase-3 activity was significantly elevated at all treatment conditions (Fig.3c, $p < 0.05$ at 0.5 mM, $p < 0.0005$ at 1 mM, and $p < 0.005$ at 1.5 mM) relative to the control group. Activity significantly increased from 0.5 mM to 1 mM and from 0.5 mM to 1.5 mM ($p < 0.05$). Therefore caspase-3 activity was not suppressed by any of the zinc concentrations evaluated.

5.3.4 Effect of transition metal ions on caspase activity

In order to investigate whether the transition metal ions of common degradable metals stimulate caspase activity at excess concentration, we performed activity assays with arteries that were incubated in 0.5 mM solutions of either Fe, Mg, or Zn. Zn elicits the strongest increase of caspase-3 activity as compared to Mg and Fe transition metal ions (Fig.5-3.D, $p < 0.0005$).

Fig.5-3 Caspase activation in the ex vivo arterial culture system. Caspase -9 (A), -8(B), and -3(C) activity in 3 different concentrations is shown. Caspase-3 activation in 500 μ M Fe, Mg, and Zn is shown in (D). E and F represent confocal images from control and Zn treatment groups respectively of active caspase 3 staining present in cross-sections. Data error is presented in sample standard error and *=p<0.05, **p<0.005, ***p<0.0005, and ****p<0.00005.

5.3.5 Addition of Cu to bulk zinc materials

A promising Zn material alloyed with copper (Cu) was selected to evaluate the potential biological effect of its eluted Cu constituent on caspase activity. Fig.5-4 shows the neointimal response to Zn-4Ag-0.6Mn, and Zn-4Ag-0.8Cu-0.6Mn-0.15Zr alloy wires implanted within the abdominal aorta for up to 6 months. The mural (Fig.5-4a) and luminal (Fig.5-4b) portions of Zn-4Ag-0.6Mn wire neointimas (outlined by the blue arrows) failed to endothelialize. In contrast, a stable endothelium is clearly seen for the Zn-4Ag-0.8Cu-0.6Mn-0.15Zr wires (Fig.5-4C and D - identified by the red arrows).

ICP analysis in Fig. 5-4E shows the elemental profile of *in vitro* corrosion fluid for the two respective alloys at 14 and 28 days. Significant Cu release can be seen for the Cu-containing alloy, with no Cu release for the Cu-free control (red line denotes detection limit for the ICP system to Cu). The ICP results demonstrate the release of Cu ions into the corrosion fluid from the Cu-containing alloy. To evaluate whether Cu could induce caspase activity, we incubated arterial segments in 5 μ M and 50 μ M Cu solutions for 12 hours and probed for caspase-3 activity. Neither condition elicited significant caspase activity within smooth muscle cells, indicative of caspase-independent mechanisms of biologic interaction.

Fig.5-4 Copper addition in advance zinc alloys effects the vascular response through non caspase dependent pathways. A and B show mural and luminal 600X normal magnification H&E images of Zn-4Ag-0.6Mn wires at 6 months within the abdominal aorta. Blue arrows depict inflamed endothelium. C and D

show luminal and mural views Zn-4Ag-0.8Cu-0.6Mn-0.15Zr wires at 6 months in histological cross-section. Red arrows show confluent, non-inflamed endothelium. E shows ICP analysis of corrosion media

5.4 Discussion

The present study explored the similarities and differences between the neointimas that develop around degradable (zinc) and non-degradable (platinum) implant metals. From a histomorphometric perspective, platinum and zinc materials placed into healthy murine arteries form neointimas with similar characteristics (Fig.5-1). From a broad view, the similar cell densities, neointimal area, and neointimal thickness measurements demonstrate the promise for use of zinc-based metals as vascular implants.

Materials selected in the past to serve as stents, either as biostable or bioresorbable materials, are intended to be highly bioinert in order to avoid interaction with local and systemic systems. Although these materials perform superbly when tested in healthy animal arteries, they often fail to prevent progressive intimal hyperplasia and restenosis when deployed into the atherogenic environment of diseased human arteries. The limitations of this approach have necessitated the incorporation of drug-eluting coatings, in an effort to impart suppressive properties to bioinert stent materials.

Here we show that zinc ions elution from metallic zinc platforms potently suppresses neo-intimal hyperplasia of smooth muscle cells (SMCs), widely recognized as the primary cause of failure for stented human arteries [21-23]. We have recently reported on the promising *in vivo* behavior of metallic zinc implants in the abdominal aorta of Sprague Dawley rats, out to 6 months [16]. Confluent endothelialization was found along the luminal surface of the stable neointima, with a decreasing gradient of smooth muscle cells (SMCs) close to the implant surface [16]. Based on these preliminary observations, we hypothesized that the byproducts of zinc corrosion may exert suppressive effects on arterial SMCs. The present findings implicating caspase enzymatic activity and apoptosis signaling pathways in mediating the suppressive effects strongly suggest that the

response to ionic zinc is programmed, rather than a response to injury. If zinc ions were toxic to SMCs, we would have observed inflammation, necrosis, and progressive intimal hyperplasia.

Zinc is a well-known inhibitor of caspase-3, 8, and 9 activity when evaluated in concentrations exceeding 100 μ M [24-26]. Although this inhibitory effect has been described extensively in literature using cell-free systems [26], the relationship between excess zinc and caspase enzymatic activity using intact cell systems is less clear. For instance, it has been shown that excess zinc at high concentrations can provoke mixed forms of cell death in cancer cell lines, mediated by caspase cascade activation [27]. However, studies focusing on the effects excess zinc in vascular smooth muscle cells is relatively lacking. Because the concentration of zinc is expected to be high directly at the interface of a zinc-based implant, we sought to measure caspase activity in neointimal cells at the interface of zinc wires implanted into the arterial environment and in arterial media layers cultured *ex vivo* in simulated high zinc environments.

Caspases 3 and 8, but not 9 become significantly elevated in high concentration zinc conditions, as shown by Fig.5-3. Surprisingly, at concentrations of zinc exceeding 0.5 mM, caspases 8 and 9 dramatically decrease in activity relative to both the control and 0.5 mM conditions. Conversely, caspase-3 continues to increase in activity with zinc concentration, consistent with the extensive apoptosis we have detected in the neointimas of zinc wires. This is also not surprising, as caspase-3 is the most robust caspase in respect to inhibition of the proteolytic activity, requiring 1 mM for complete inhibition of the enzyme [26]. Caspase-9 is known to be potently inhibited by zinc [24], which could explain its lack of activation in the lower concentration condition and suppressed activity at the higher zinc levels. Caspase-8 may also be inhibited in some manner at high concentrations of ionic zinc. The data suggests a complex interplay of caspase activation and zinc concentration, which ultimately yields caspase-3 activation and apoptosis execution.

We also found an anti-inflammatory effect from the added elution of copper ions, which significantly reduced neointimal thickness and inflammation around the implant. This effect is independent of the caspase-3 induced activation of zinc, as shown in Fig.5-4f. It has been reported that both Zn^{2+} and Cu^{2+} can stimulate NO release from endogenous and synthetic S-nitrosothiols (RSNOs) [28-30]. We recently reported that the amount of NO released by Cu^{2+} from the synthetic RSNO, SNAP, was an order of magnitude higher than that of Zn^{2+} [31]. Free endogenous RSNOs circulate in the blood and are constantly replenishing [32], providing a continuous source of NO generation for degradable materials that elute Zn^{2+} and/or Cu^{2+} . Local NO production from the release of Cu^{2+} from biodegradable materials in addition to Zn^{2+} may improve the biological response, owing to the critical physiological roles of NO in regulating vascular function. Beneficial effects of NO include inhibition of platelet aggregation [33], leukocyte adhesion [34, 35], and vascular SMC proliferation [36], all contributing factors to the restenosis of stented arteries. Interestingly, it has been shown in peripheral blood mononuclear cells that zinc protects against copper induced DNA damage, and substantially raises the LD50 value of copper [37]. Copper also increases the proliferation rate of endothelial cells, but not SMC's [38]. Fig.5-4 nicely shows a confluent, endothelium that is largely lacking in the inflammatory ridden ternary alloy.

The present findings have major implications for the future of bioresorbable stent development. Drug eluting polymer coated stents (DES) have become the main approach for the revascularization of occluded coronary arteries. These stents combine mechanical scaffolding with the release of pharmaceutical agents from a polymer coating on the stent surface that act to inhibit smooth muscle cell proliferation [39]. Although second generation DES have reduced the rates of late stage thrombosis relative to bare metal stents (BMS) [39], these stents still experience a ~10% incidence of in-stent restenosis (ISR) [40]. Furthermore, ISR for DES begins after 6 – 9 months and increases up to 2 years following implantation, as opposed to peaking between 3 – 6 months for BMS [40, 41]. The time course suggests limitations related to the drug in addition to the inherent limitations of permanent stents. For instance, the drug release time course is limited to

several months. Of potentially major importance when considering drug safety, paclitaxel has recently been shown to sharply increase the risk of death many years after stent deployment, likely due to systemic side effects [42]. It is an open question whether other commonly used drugs for stent elution elicit similar systemic effects.

Recognized limitations with DESs have given rise to the concept of fully bioresorbable polymeric scaffolds (BRS), which are in ongoing development to replace their permanent metallic counterparts [43]. This feature is expected to restore positive arterial remodeling while removing sources of thrombogenesis, inflammation, and neointimal activation. However, all the polymers that have been selected for scaffolding exhibit substantially reduced mechanical properties relative to metals, necessitating thicker struts. This worsens malapposition, reduces endothelial regeneration, and leads to an increased risk of thrombosis [43]. Bioresorbable metallic alternatives are further behind in development, yet with appropriate modification can achieve similar mechanical properties as conventional stent metals and have achieved some preliminary success [43]. Unfortunately, Fe and Mg based scaffolds will require drug eluting polymer coatings when deployed into small diameter arteries.

Due to the limitations of anti-proliferative eluting drugs, the development of a bioresorbable metallic stent that does not require synthetic drug elution is urgently needed. In marked contrast to what has been shown previously for Fe and Mg, we have shown here that the byproducts of zinc implant biocorrosion suppress the harmful activities of neo-intimal smooth muscle cells. Zn^{2+} and Cu^{2+} , like Mg^{2+} and Fe^{2+} , are present in the body naturally and the low quantities released from a stent that biodegrades over a year, in particular with copper included as a low concentration alloying addition, are unlikely to substantially alter their systemic levels. Thus, Zn^{2+} and Cu^{2+} may act in a similar fashion as synthetic eluting drugs to protect against intimal hyperplasia of smooth muscle cells, yet for a longer duration and without the harmful systemic side effects of synthetic drugs. We propose, consequently, that the development of bioresorbable metallic zinc stents could proceed without polymer coatings or synthetic drug elution.

5.5 Conclusions

In the present paradigm, byproducts of implant degradation are seen largely as systemic pollutants to be cleansed from the body without promoting toxic effects. We propose a new paradigm, wherein the appropriate degradation byproducts are seen as therapeutic agents, similar to the synthetic drugs impregnated into modern day drug-eluting stents, yet naturally found in the body. Due to the physiological presence of the element, an elevated local concentration can be achieved to deliver therapy near the implant without promoting systemic effects. It is straightforward to embed multiple components within the zinc matrix and control their concentrations through conventional alloying approaches. Such an approach would serve both to improve the structural and mechanical properties of the material and to elute multiple therapeutic agents into the local environment.

5.6 Acknowledgments

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6 Concluding Remarks

This work illuminates the complex interplay between degrading zinc-based implant materials and the vascular environment. Current simplistic conceptualizations of the foreign body response must be reconsidered when examining degradable materials. The field of biomaterials has witnessed a fevered introduction of degradable materials over the past 20 years, including metals, polymers, composites, additively manufactured matrices, and natural substrates. With the addition of zinc as a new class of biomaterials, our understanding of how the innate and adaptive immune systems interact with implanted materials must evolve. In the body of evidence provided in chapter 1, we have shown that the inflammatory response is capable of accommodating degrading zinc materials. Identifying links between cellular driving forces that increase corrosion activity and vice versa could allow for formulating a theoretical foundation that further enhances our abilities to engineer advanced degradable biomaterials.

When designing zinc-based biomaterials for cardiovascular use, it is imperative to observe the material in the “application environment”. Conventional 2-D cell culture methodology does not provide meaningful insight when applied to materials intended to operate in the complex environment of the intraluminal arterial space. Conversely, manufacturing of stents and implantation in pig models is prohibitively expensive. To bridge the gap, we developed a simple and effective methodology to accelerate the invention of novel degradable biomaterials and accomplish their screening based on their quantified performance in a realistic experimental rat model. Using this approach, presented in detail in chapter 2, material characteristics in a wire implant geometry can be described in terms of their contribution to neointimal growth. To explore this approach, the author evaluated different surfaces with varied surface oxide layer stability engineered on zinc substrates, presented in chapter 3. The reduction in biocompatibility for one of the surfaces was related to a relatively low surface oxide stability that increased susceptibility for pitting corrosion. We found that pitting corrosion vs. uniform corrosion leads to localized bursts of zinc corrosion, producing in turn a highly localized inflammatory reaction that worsens neointimal growth. This led us to conceptualize the

engineering of zinc implants with a “tunable” neointimal response by controlling the surface oxide film stability. Taken together, the chapter illustrates how relating material characteristics to the micro cellular and macro tissue development provides fundamental insight into the neointimal responses, which in turn can impact the metallurgical engineering of the material.

In addition to exploring relationships between material properties and biological responses, the framework of testing that we developed in chapter 2 can be applied to the purpose of identifying the best performing materials fabricated by our research group. By statistical comparisons of biocompatibility metrics using high sample sizes, we have identified the quinary alloy (Zn-4Ag-0.8Cu-0.6Mn-0.15Zr) as the best performing material, following solution treatment to maximize mechanical properties. This is the first alloyed zinc material that performs better than pure zinc. The identification of this alloy is described in detail in chapter 4. The mechanisms by which this material generates outstanding biological responses is clarified in chapter 5 and summarized as follows:

I - the addition of Cu reduces inflammation and increases endothelialization

II - the provocation of inflammation by AgZn₃ precipitates is decreased by solution treatment

III - zinc and copper exert a synergistic effect to suppress neointimal growth

In chapter 5, the author begins to clarify the mechanisms by which zinc based degradable materials may suppress neointimal growth. When the neointimal development around platinum and zinc wires was compared, there was a reduced smooth muscle cell presence and an increased cellular apoptosis in the zinc neointima. The author hypothesized that the release of Zn²⁺ from degrading zinc-based implants induces programmed cell death, or apoptosis, of smooth muscle cells. The role of Zn²⁺ was confirmed by exposing arterial smooth muscle cells to high concentrations of Zn²⁺, ex vivo, in order to isolate the effects of ionic zinc from variables related to implant materials. The activity of the executioner caspase (3) was found to be elevated in high zinc conditions. This controlled form of cell

death is beneficial to a progressively developing neointima, as it would shift the death-proliferation balance to stabilize against the progressive growth that eventually occludes stented arteries. This is an extremely important discovery since the proliferation of smooth muscle cells represents the hallmark mode of failure for arterial stents.

In summation, the work begins with the development of a new methodology to qualitatively and quantitatively evaluate the in vivo biocompatibility of degradable implant metals. This methodology was used to relate material characteristics to biological responses, including surface oxide film character, alloying composition, and bulk processing conditions. In addition to generating insights into fundamental relationships between materials and biocompatibility, the approach can be used to rank materials in terms of biocompatibility. From this approach, beneficial alloying elements and processing conditions can be identified. Beneficial elements identified by these approaches can be further investigated for their therapeutic value in the form of transition metals, since all the elements in the implant will be released due to biocorrosive activity.

Future studies will need to explore important questions that were generated by the present work. The present work evaluates materials in a normal, healthy animal system. While necessary to clarify fundamental relationships between material characteristics and neointimal formation and begin to clarify mechanisms of suppression by ionic zinc, a diseased arterial environment is more representative of the clinical environment. In contrast to a healthy artery, the diseased atherosclerotic environment is pro-inflammatory, more complex and unique per individual, and lipid laden. The performance of degradable zinc materials will need to be evaluated in a dynamic and pro-inflammatory micro-environment. The most suitable modeling systems for arterial disease have been developed decades ago by molecular biologists, using transgenic mice. It should be possible to adapt the implant surgery and our quantitative morphometric approach to the mouse. Positive performance of degradable zinc-based materials in diseased arteries would increase confidence in the successful clinical translation of materials selected using the approaches described in the present dissertation.

A Appendix for Chapter 1

Fig. 1-S1. Quantification of CD163 staining using average pixel counting of multiple tissue sections per sample.

Fig. 1-S2. Quantification of CD11b/c staining using average pixel counting of multiple tissue sections per sample.

Fig. 1-S3. In depth clarification for Fig. 2. Yellow arrows in panel A show mononuclear cells. Cyan asterisk in C show region of compact corrosion product with no nuclear staining. Red asterisk in D depict region of necrosis. Scale bar is set to 100 μ m.

Fig. 1-S4. Second clarification of Fig. 2. Red asterisk in E show necrotic regions, yellow arrow displays a representative normal nuclear morphology, and the green arrows depict representative abnormal nuclear morphology. Red arrows in F identify thick, circumferentially aligned collagenous fibers, while in panels G and H show a distinct collagenous capsule. Blue asterisks in panel G identify chronic inflammation outside of the capsule, while in panel H they identify the chronic inflammation both inside and outside the capsule. Scale bar is set at 100 μ m.

Fig. 1-S5. Red asterisks for the SHG explant at 1.5 and 3 months show large amounts of corrosion product. Yellow arrows show representative viable mononuclear cells. The red line for the bottom two panels depict the original border of corrosion pocket, with the red arrows showing the direction of cellular penetration. Light blue arrow in bottom right panel shows a representative region of pyknosis and karyorrhexis. Scale bar is at 100 μ m

Fig. 1-S6. Yellow arrows in top left panel show representative mononuclear cells, while the red arrows in the top right panel portray the cellular movement from the original tissue - corrosion product border inwards towards the corroding implant. The green arrows in the bottom panels show neovascularization, and the cyan arrows identify representative fibroblasts. The yellow asterisks in the 2AI 4.5month panel shows corrosion product impregnated with cell nuclei. The red circles in the bottom right panel show sites of macrophage fusion. Scale bar is set at 100 μ m.

Fig. 1-S7. Cyan arrows in the 4-AI 3 month panel show representative mature fibroblasts at the interface. Red asterisks show representative inflammatory cells. Green arrows in the bottom right panel show neovascularization. Yellow arrows depict aligning fibroblasts. Scale bar is set at 100 μ m.

Fig. 1-S8. Yellow arrows in the top left panel show acellular collagenous bundles, with the red circle identifying representative dying inflammatory cells within the corrosion layer. The 4-Al 6 month panel's red asterisks show a thick, acellular collagenous capsule, with the red oval identifying inflammatory cells. The cyan arrows in the Zn-8Al 3.5 and 6 month panels identify the fibrous capsule. The scale bar is set at 100 μ m.

Fig. 1-S9. The top left panel's yellow arrows identify representative large mononuclear cells. In the 1.5 month SHG panel, the green arrows identify representative intact cells, while the red circles identify characteristic piknosis and karyorrhexis. The green arrows in the 6 month 4N panel also show viable cells, while the red circles show areas of cell death. In the 6 month SHG panel, the white oval shows a region incorporating both viable and non-viable nuclei (green and red arrows respectively). Scale bar is set to 100 μ m.

Fig. 1-S10. Yellow arrows in 1.5-month 2-AI panel identify cells of normal nuclear morphology. Green circles of all panels show highly irregular nucleic features with increased DAPI strength. Red circles show areas void of cell nuclei. Scale bar is set at 100 μ m

B Appendix for Chapter 3

Table 3-S1. Fitting parameters of EP and AD samples after different immersion times

	Time	R_{sol} (ohm·cm ²)	Q_i (μF)	η	R_i (ohm·cm ²)	Q_o (μF)	η_o	R_o (ohm·cm ²)
EP	1	15.64	0.34	0.81	176	177.34	0.78	270
	24	14.54	1.12	0.73	40.54	268.10	0.97	180
	72	15.93	1.59	0.95	120.8	238.15	0.75	324
	120	17.7	8.75	0.95	165	920	0.75	374
	168	16.48	7.09	0.78	139	854	0.92	305
	Time	R_{sol} (ohm·cm ²)	Q_{dl} (μF)	η	R_{dl} (ohm·cm ²)	Q_{ct} (μF)	η_{ct}	R_{ct} (ohm·cm ²)
AD	1h	15.74	62.21	0.98	168.58	110.78	0.81	129.40
	24h	14.88	51.25	0.83	192.30	90.20	0.61	190.80
	72h	12.55	51.4	0.77	471.36	95.05	0.92	245.10
	120h	12.08	51.50	0.84	578.60	52.04	0.93	533.30
	168h	12.48	45.74	0.95	349.02	38.31	0.81	788.20

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Author: Roger J. Guillory, Alexander A. Oliver, Emma K. Davis et al

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Author: Roger J. Guillory, Malgorzata Sikora-Jasinska, Jaroslaw W. Drelich, et al

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