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EFFECT OF NUMBER OF CROWNS ON THE CRUSH RESISTANCE IN OPEN-CELL STENT DESIGN

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Our group is developing novel preferential covered carotid stents aimed at preventing friable parts from atherosclerotic plaques from dislodging into the cerebral blood flow, and also preserving external carotid artery (ECA) perfusion through slits on the membrane. Enhanced ECA flow rates can be achieved by designing bare metal stents with larger cells that can accommodate more slits for enhanced blood flow to the ECA where the stents have fewer crowns in the circumferential direction. The general perception is that the stent stiffness and thereby the crush resistance will decrease with fewer crowns in the circumferential direction. However, we observed the opposite effect. To study the effect of crown number on crush resistance of stents, finite element analysis (FEA) was used to evaluate the crush resistance of open-cell stent designs by varying the number of crowns. From FEA simulation results, it was found that the crush resistance of the open-cell stent design actually increases with fewer crowns. To verify this effect, three stent designs with different crown numbers were fabricated and subjected to crush resistance experimental testing. The experimental testing further confirmed the effect observed by FEA. Finally, a simplified analytical model was proposed to explain why the crush resistance of stent increases with a reduction in the number of stent crowns. From this study, we can infer that the stent's crush resistance increases with reduction in the number of stent crowns in the circumferential direction.

1. Introduction

Our group [Kabinejadian et al. 2013; 2015a; 2015b; Kumar et al. 2016] has been actively developing novel covered stents for preventing small friable parts of atherosclerotic plaques from dislodging into the cerebral blood circulation, and at the same time, preserving external carotid artery (ECA) blood flow. The device comprises of a bare metal stent coated with a biocompatible polymer membrane which has several arrays of cut slits. The main purpose of the slits on the covered membrane was to let blood through the membrane into ECA while confining the plaques along vessel wall and prevent them from dislodging. In one of our earlier in-vitro investigations [Kabinejadian et al. 2015a] we showed that our covered stent design is more efficient in emboli prevention than its corresponding bare metal stent, and at the same time, preserves more than 83% of the original flow of the ECA in carotid artery bifurcation models. Our group evaluated this covered stent design [Kabinejadian et al. 2013; 2015b] (with slit openings on ePTFE membrane) in various models of arteries with different arterial diameters, curvatures, and side-branch angles. We aimed to design new bare metal stents with bigger cell size which can provide higher ECA flow rates through enhanced perfusion to the side-branch as they can accommodate higher number of slits. Larger stent cell size thereby contributes to enhanced slit distribution on the membrane (and

Keywords: stent crown, nitinol, membrane, carotid, stent crush resistance finite element analysis.

thereby better side-branch flow preservation) and desirable crimpability [Kumar and Cui 2016; Kumar et al. 2014]. As part of our stent design-iteration process, we reduced the number of crowns of our open-cell stents in the circumferential direction to get bigger cells and put them through finite element analysis (FEA) to evaluate their crush resistance. Crush resistance can be defined as the ability of a stent to withstand crush loading and therefore prevention of stent collapse is crucial for stent development especially for superficial locations like carotid arteries [García et al. 2012; Timaran et al. 2011; Duerig and Wholey 2002] and peripheral indications as per Food and Drug Administration (FDA) standards [Dhruva et al. 2009; US Food and Drug Administration 2010]. The aim of this study is to evaluate the effect of the number of stent crowns in the circumferential direction on the stent crush resistance through the following steps: (a) analysis of this effect by FEA based modeling and simulation for three different stent designs; (b) fabrication of the stent designs followed by experimental compression tests; (c) providing some explanations with simplified analytical study.

2. Materials & methods

2.1. Stent design. In this study we took reference from the commercial stent E-Luminexx[®] stent (Bard Peripheral Vascular Inc., Tempe, AZ) to model the stent which has 12 crowns in the circumferential direction. Additionally, we designed two more stents by reducing their crown numbers to 9 and 6, respectively in the circumferential direction. Figure 1 shows the models of all the three stents. SOLIDWORKS (Dassault Systemes, MA) was used to model the stent geometries and ABAQUS (Dassault Systemes, MA) to conduct FEA. The stents had an outer diameter of 6 mm, a length of 40 mm and struts with uniform thickness of 0.15 mm. These are hybrid stents with combined [Timaran et al. 2011] open- and closed-cell sections (Figure 3). The open-cell sections improve flexibility and enhance stent-vessel conformability without injuring the vessel [Pierce et al. 2009; Carnelli et al. 2011; Tadros et al. 2012]. The closed-cell sections provide sufficient radial strength and in the case of lesions, to secure plaques and hold the plaque in position [Tadros et al. 2012; Hart et al. 2006; Bosiers et al. 2007]. The specifications of stent models considered in this study are given in Table 1. All the three stents were laser-cut from a nitinol tubes of 2.0 mm outer diameter (OD) and 0.18 mm thickness. They were then expanded and shape set using a cylindrical mandrel to 6 mm $\times$ 40 mm (OD $\times$ length) following which electropolishing was done to get a final strut thickness of 0.15 mm. The fabricated nitinol stents for testing are shown in Figure 4.

2.2. Material model. Nitinol has been used as a stent material as it possesses excellent properties like superelasticity, shape memory, corrosion resistance, biocompatibility, radiopacity, and fatigue resistance [Hansen 2008; Duerig et al. 1999]. When cooled to a very low temperature, nitinol transforms fully to martensite and becomes easily deformable and hence can be effortlessly crimped into a small catheter. When the stent is released into the artery, it recovers to its predetermined, original diameter when exposed to the body temperature which is higher than the Austenite finish (Af) temperature. The built-in Nitinol UMAT was used for our stent simulations. In this model [Auricchio and Taylor 1997; Auricchio et al. 1997] the total strain is the sum of the elastic, transformation and the plastic strain. The mechanical properties of nitinol used in this work are partly from our previous work [Kumar et al. 2013; 2014] and partly based on the information given by our stent fabrication partner and are shown in Table 2.

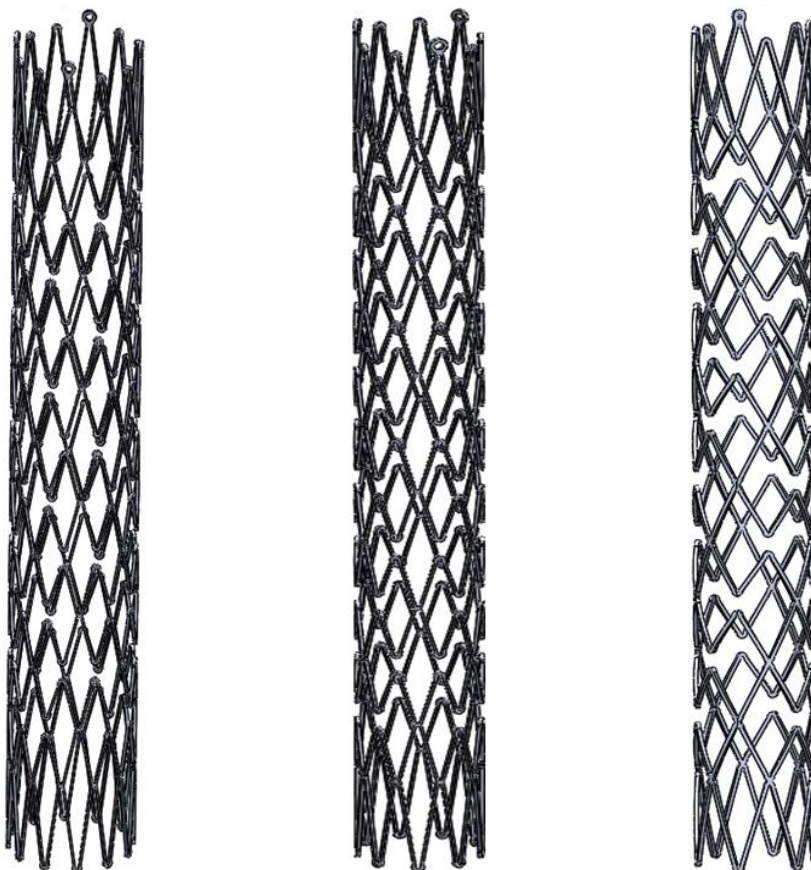


Figure 1. Stent designs investigated in the study: 12 crown design (left), 9 crown design (middle) and 6 crown design (right).

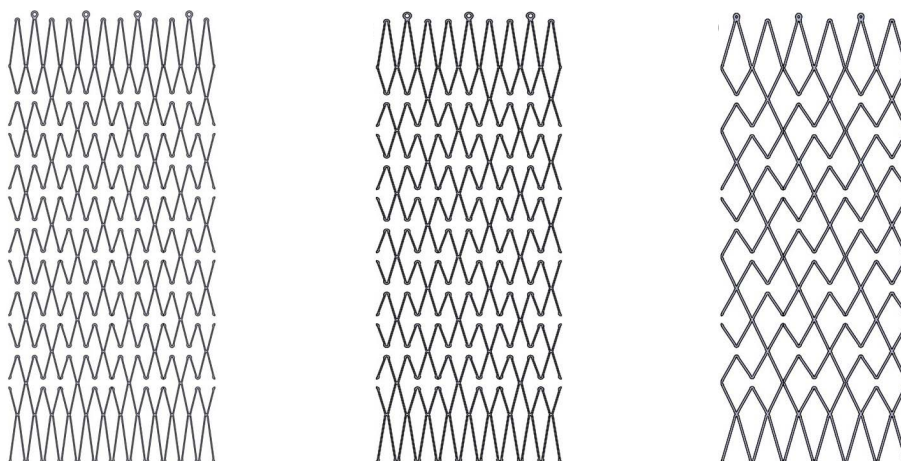


Figure 2. Stent designs with different crown numbers shown in their planar form: 12 crown design (left), 9 crown design (middle) and 6 crown design (right).

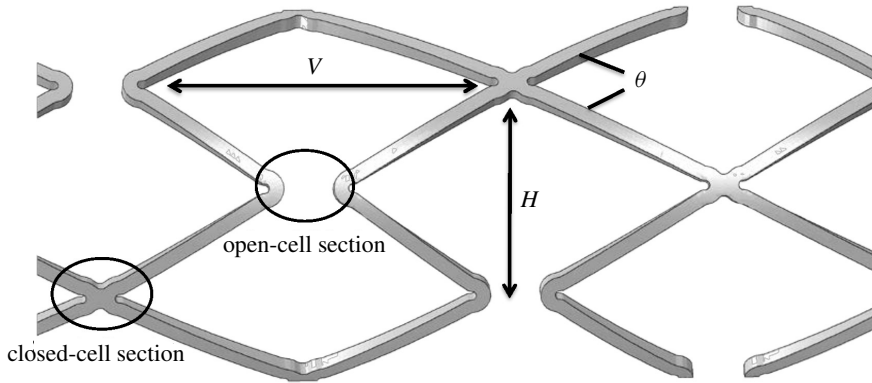


Figure 3. Representation of the stent cell specifications of the stents used in this work (V stands for the distance in axial direction, H the distance in circumferential direction and θ the inter-strut angle).



Figure 4. The three fabricated stents with varying crown numbers in the circumferential direction: 12 crown design (left), 9 crown design (middle) and 6 crown design (right).

stent	12 crown design	9 crown design	6 crown design
outer diameter, OD (mm)	6	6	6
length (mm)	40	40	40
number of crowns	12	9	6
vertical inter-strut distance, V (mm)	5	5	5
inter-crown distance, H (mm)	1.5	2.25	3
inter-strut angle, θ (degree)	21	33	45
strut thickness (mm)	0.15	0.15	0.15
strut width (mm)	0.15	0.15	0.15

Table 1. Specifications of stent models considered in this study.

variables	nitinol
Austenite elasticity, E_A (MPa)	40000
Austenite Poisson's ratio, ν_A	0.33
Martensite elasticity, E_M (MPa)	32000
Martensite Poisson's ratio, ν_M	0.33
transformation strain, ε^l	0.041
loading $(\delta\sigma/\delta T)_l$ (MPa T^{-1})	6.7
start of transformation loading, σ_l^s (MPa)	465
end of transformation loading, σ_l^e (MPa)	480
reference temperature, T_0 ($^{\circ}C$)	22
unloading, $(\delta\sigma/\delta T)_u$	6.7
start of transformation unloading, σ_u^s (MPa)	260
end of transformation unloading, σ_u^e (MPa)	248
start of transformation stress in compression, σ_{cl}^s (MPa)	-449
volumetric transformation strain, ε_v^l	0.041
Af temperature ($^{\circ}C$)	37
strain limit, ε_{max}	10%

Table 2. Nitinol material properties.

2.3. Computational simulation. The ABAQUS/Standard (v. 2017) FEA package was used to carry out the nonlinear FE analysis. The stents were modeled with eight-node linear brick elements with reduced integration and hourglass control. To make sure all the results were independent of further mesh refinements, mesh sensitivity studies were conducted. Mesh convergence was achieved by sequentially increasing the number of elements until there was no appreciable difference in the stress at the end of the analysis. The final mesh of the 6 crown, 9 crown and 12 crown stents comprised of 43,373, 72,481 and 103,845 elements, respectively. For the flat plate compression test, similar to an earlier work [García et al. 2012], each stent was placed between two rigid plates separated by a distance of 6.5 mm. The stents were then compressed by imposing a displacement of 3 mm to the upper plate, similar to the experimental test (described in Section 2.4), while constraining the lower plate in all degrees of freedom

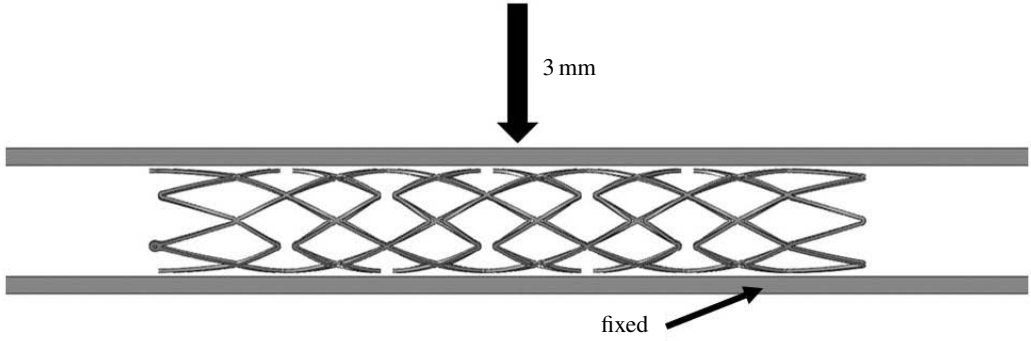


Figure 5. Boundary conditions used in the simulation for radial compression. The lower plate is fixed in to prevent rigid body translation in all three directions.

(see Figure 5). Contact between the plates and the stents was modeled with frictionless tangential and hard normal contact. A penalty interaction property was used to enforce impermeable boundaries. The reaction force after full loading was recorded.

2.4. Experimental setup. All tests were performed using an Instron 3345 machine (Instron, 825 University Avenue, Norwood, MA 02062, USA) at 20 °C–22 °C. For radial compression test which is similar to the experimental test [Petrini et al. 2005] developed to estimate the radial force of stents, the loads were applied over the full length of the stents with a plate 50 mm in diameter. All stents were placed between two flat plates. No external supports were used for the stents. Three samples were tested for each design (6, 9 and 12 crowns). For the individual stents, the test was repeated three times rotating the stent twice by 120° to negate the possible influence of the rotational orientation of stent. A displacement rate of 20 mm/min was used to start the test until a displacement of 3 mm was achieved. The force and displacement data were recorded using the Instron Bluehill 3 modular applications software for both loading and unloading processes. The experimental setup for the radial compression test is shown in Figure 6.

3. Results

3.1. Simulation. The results obtained from simulation for the three flat plate compression tests are presented in Figure 7. The results show that the force required to produce a displacement of 3 mm increases with a reduction in the number of crowns. These forces are 5.13 N, 1.03 N and 0.77 N for 6 crown, 9 crown and 12 crown stents, respectively. From the results it can be inferred that the 6 crown stent shows higher crush resistance among the three designs in comparison to the other designs whose crown number increases.

3.2. Experiment. Response to experimental compression tests is presented in Figure 8. The curves in the experimental results plots represent the mean of the three individual test data for each design. From the experiment, it is evident that the crush resistance of the 6 crown stent is the highest followed by the 9 crown and 12 crown stents. This trend is in agreement with the inference from our simulation results.

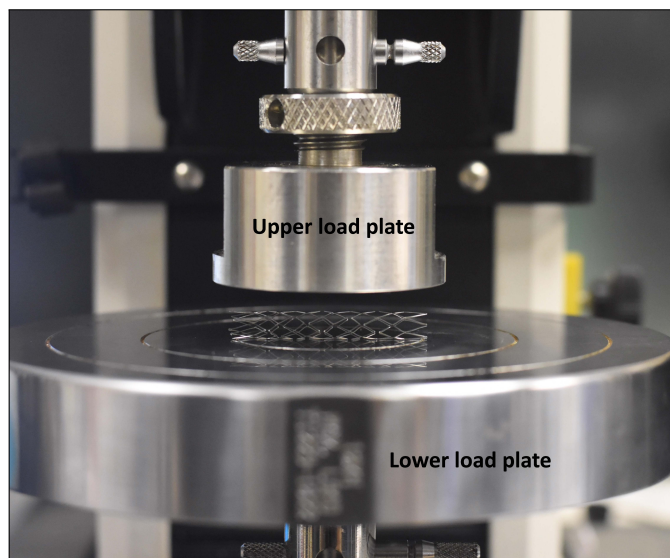


Figure 6. Experimental setup for radial compression test.

4. Discussion

From the simulation and experimental results it could be inferred that even though the metal density reduces as the crown number is reduced in the circumferential direction, surprisingly and contrary to the common observation, the crush resistance of stents increases. There could be a couple of reasons for this

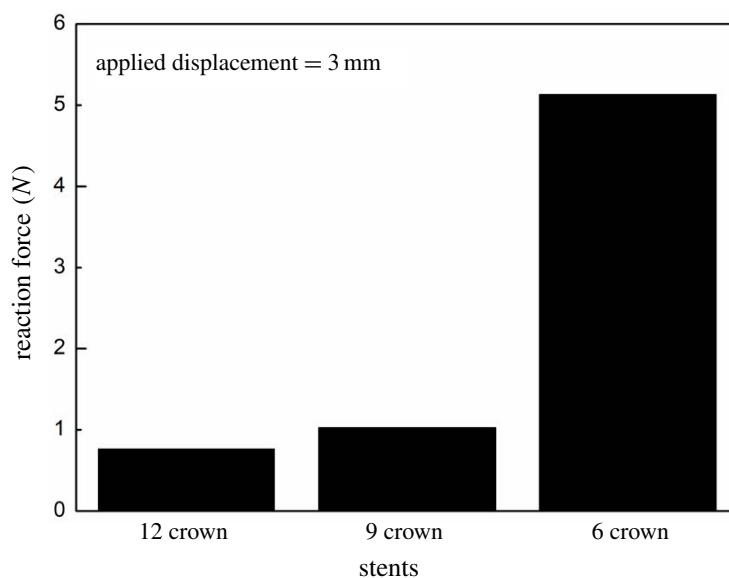


Figure 7. Results of the FEA study: 6 crown stent having maximum reaction force among the three designs in agreement with our experimental results.

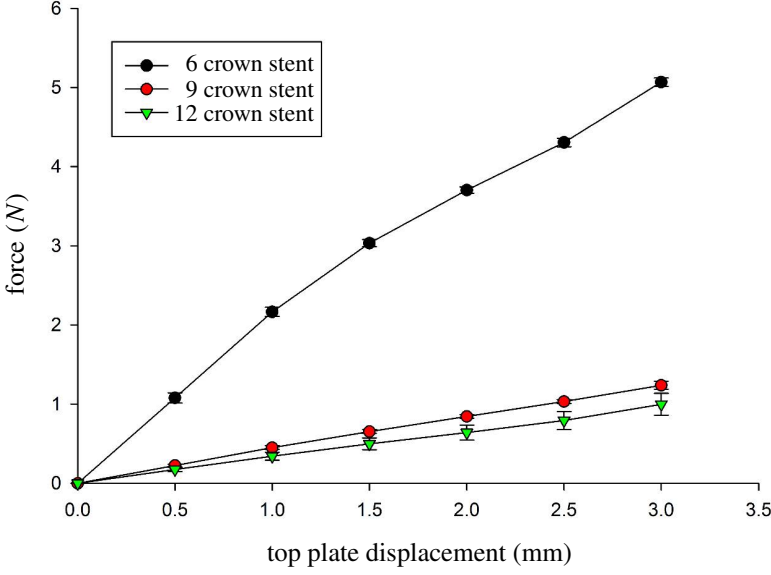


Figure 8. Results of the Experimental study. Force-displacement curve for the radial compression loading of the three stents.

trend. Firstly, when the number of crowns keeps getting reduced in the circumferential direction, there is also a proportional reduction in the open-cell sections which causes the stent to have high stiffness [Carnelli et al. 2011; Moore 2012; Hijazi et al. 2013] as open-cells offer lower stiffness. Secondly, the relationship between crown angle and the crush resistance contributing to the above trend has been explored below in details.

Each stent cell is made up of three basic elements: strut, link, and crown [Alaimo et al. 2017] responsible for crush resistance and crimpability. To understand the increase in the crush resistance with a reduction in the number of crowns, we can consider the stent crown as a simple ring as shown in Figure 9. The zoomed-in part shows one-half of the crown that is considered as a simple beam for the below analytical model.

Let H be the ring height, L the total length, n the number of rings, $\theta = (\text{crown angle})/2$. To obtain the stiffness, a force F is applied at two ends and thereby we have

$$\Delta D = \sum_{i=1}^{2n} \frac{F \cdot \cos \theta}{k_b} \cdot \cos \theta = \frac{F}{k_b} 2n \cos^2 \theta, \quad (1)$$

where ΔD is the total displacement in the direction of the force F , and k_b is the beam stiffness. We can write the beam stiffness as

$$k_b = \frac{3EI}{l^3} = \frac{3EI}{(H/\cos \theta)^3},$$

where EI is the bending rigidity of stent strut. Thus the total stiffness k_{total} is

$$k_{\text{total}} = F/\Delta D = C(\cos \theta/n), \quad (2)$$

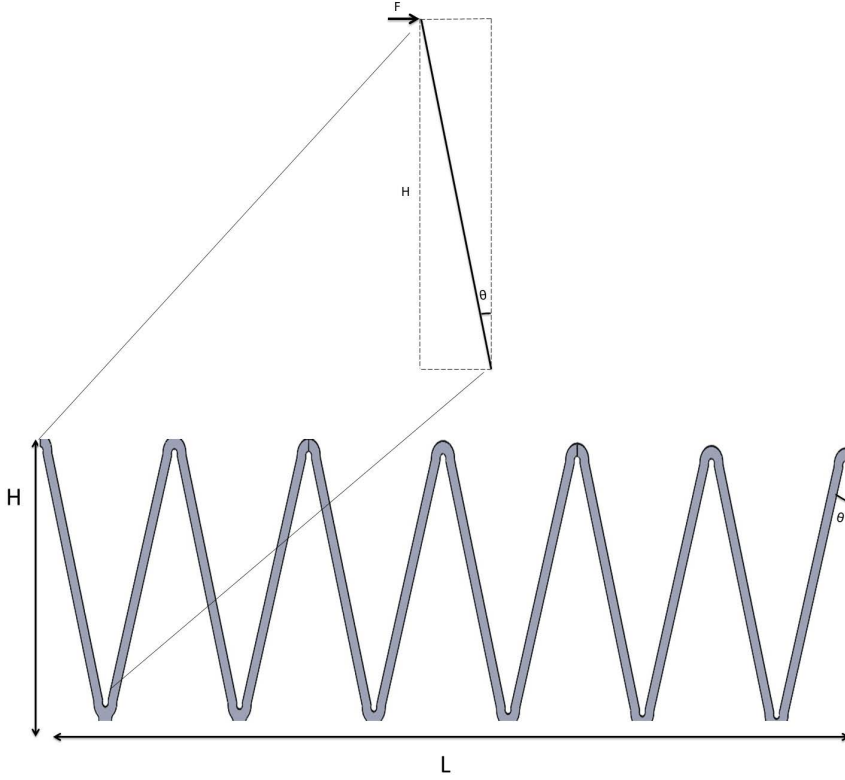


Figure 9. Stent cells being considered as a simple ring to study its stiffness behavior in relation to the number of crowns.

where $C = 3EI/2H^3$. The stiffness can be further expressed as

$$k_{\text{total}} = C \cdot \frac{1}{\sqrt{1 + (L/2nH)^2}} \cdot \frac{1}{n} = C \frac{1}{\sqrt{n^2 + L^2/4H^2}}. \quad (3)$$

Equation (3) shows that when the number of crowns increases, the crush resistance decreases due to a reduction in the total stiffness. When the number of crowns increases, the deformation per crown decreases for a given deformation, thus the load to produce the given total deformation reduces as illustrated in (3). The value of k_{total} as a function of n is plotted in Figure 10, which directly illustrates the relationship between the total stiffness (which determines the crush resistance) and the number of crowns.

5. Conclusion

In this paper the effect of number of crowns on the crush resistance of open-cell stents was studied using FEA and validated through experiment and analytical model. The conclusion made from the study is that the number of crowns in the circumferential direction was inversely proportional to crush resistance of open-cell stents. This is of particular importance especially when stents are designed for superficial locations like the carotid artery where high crush resistance is desirable due to the possibility of the application of external loads. So designing stents with lesser number of open-cells will render

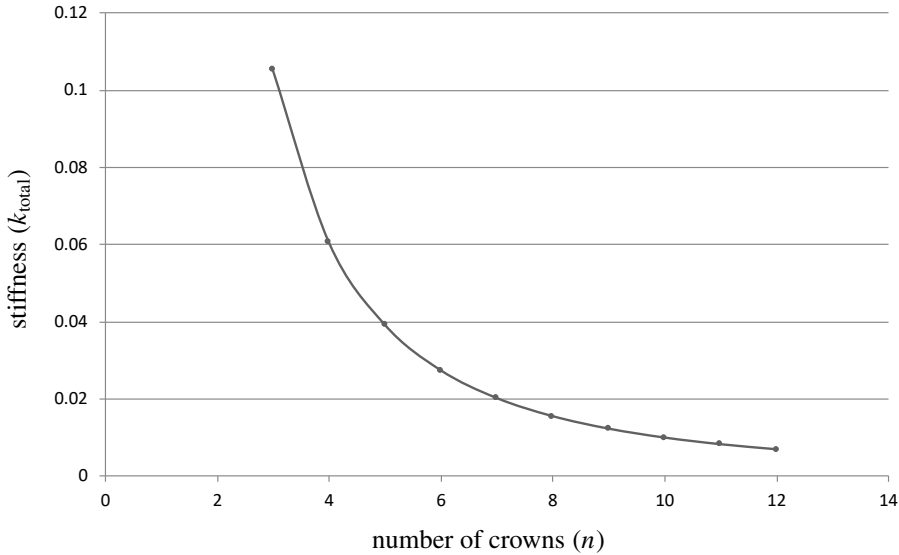


Figure 10. Total stiffness vs number of crowns with the value of k_{total} as a function of n according to (3).

them suitable for application in superficial blood vessels. This work will provide a guideline for stent design in different applications, especially at superficial locations such as carotid and femoral arteries and peripheral indications as per FDA standards where the stents are usually subjected to external loadings during the human activities of daily living.

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