A critical appraisal to the decision by the company Zimmer Biomet to withdraw the Maestro™ Wrist Reconstructive System from the marketplace

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Abbreviations

Note by the author: 75% of 104 Scientific References related to TWA are not older than 10 years, and 81% of them are not older than 5 years.

Case presentation
This is to inform the community of all Orthopaedic and Hand Surgeons around the world who are interested in Total Wrist Arthroplasty (TWA) about the official decision by the European and German Branches (Winterthur, Switzerland and Freiburg i. Breisgau, Germany) of the company for distribution (Zimmer Biomet Holdings Inc., Warsaw, Indiana/USA) for the Maestro™ Wrist Reconstructive System (WRS, insertion of the carpal plate with variable or fixed locking screws) (Figure 1A-C). Recently, the company informed the author on February 26., 2018 that they have definitively withdrawn the Maestro™ WRS from the marketplace here in Germany, and this decision will be extended in 2018 for all other countries worldwide if the CE certification is finished in these countries in 2018 as well. The company officially wrote that the cause for its withdrawal is that as follows: The Maestro™ WRS is approved only for its cemented use by the US Food and Drug Administration (FDA) and there are no publications in the literature worldwide on favorable results with its cemented use, and so, it cannot be guaranteed the surveillance of this implant by the company if it inserted in a non-cemented manner (i.e. "off-label" use).

Generally, TWA with the use of the third generation types (biaxial-anatomical, unconstrained, ellipsoid metal-on-polyethylene (PE) surface articulation) is the recognized motion-preserving treatment option to a Partial or Total Wrist Fusion (PWF or TWF) in cases of rheumatic or non-rheumatic Wrist Joint Osteoarthritis (WJOA) worldwide [1-13]. Wrist joint motion modulates the synergistic and antagonistic extrinsic / intrinsic motion at the finger joints which is an important prerequisite to realize all powerful functional grips of the hand. PWFs crossing the midcarpal or radiocarpal joints always result in loss of the overall wrist joint motion of approximately 50% [14,15]. Adams et al. demonstrated impressively in 2003 that in healthy subjects limited wrist joint motion inevitably leads to a significant worsening of their ratings in activities of daily living (perineal hygiene, buttoning a blouse or shirt, threading a needle, turning over the pages of a newspaper, opening a door with the door handle, picking up small objects, and so on), and Nydick et al. reported in 2013 that patients who received primarily a TWA for treatment of post-traumatic WJOA rated their functional outcome in Patient-rated wrist evaluation (PRWE) statistically significant better than patients who received primarily a TWF [16,17]. Adey et al. reported in 2005 that nearly all patients who received primarily a TWF would elect to have a procedure that could make their wrist move again if one were available, and Ekroth et al. reported in 2012 that their patients with failed TWA's utilizing older generation types would have a TWA again despite that their long-term results being poor and many of them being revised to a TWF [18,19]. It is still widely known from the literature that for the wrist 5° to 40° of flexion, 30° to 40° of extension, and 10°/15° of radial/ulnar deviation or 40° of combined radial-ulnar deviation are required to realize the most essential activities of daily living [20,21]. When patients are unsatisfied with their primarily performed motion-restricting TWF, then a secondary conversion to a motion-preserving TWA with a third-generation type, also utilizing the Maestro™ WRS, can be an useful option to improve the patient’s disability [22,23]. Additionally, the Maestro™ WRS has proven to be useful as well as salvage procedure of a failed previously performed PWF, and in single cases as early treatment option for highly comminuted distal radius fracture in selected older and elderly patients [24,25]. Furthermore, recent evidence suggests that the complication rate of TWA does not differ statistically significant in comparison to patients undergoing a TWF [26,27].

Till 2014, implant survival with the third TWA generation types was reported to be 90 to 100% at 5 years in most series, but it declines from 5 to 8 years [9]. These results are completely comparable with those after Total Shoulder / Elbow / Ankle Arthroplasties which are much more less debate in the literature than TWA. Recently, rheumatoid arthritis remains the most common indication for TWA with a relative portion ranging from 51 to 71%, and followed by 14% for treatment of post-traumatic WJOA of all TWAs performed by surgeons who have published their experiences with this procedure [9,26]. Two questions are not clearly answered currently: first, how can patients load their wrists with a TWA; and second, are young patients suitable receiving

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assessed with good bone stock (including the Maestro™ WRS), and towards the increasing expansion of the indication also for selected young and active [22-32]. However, there is a trend in the literature lifestyle, whereas patients with post-traumatic WJOA are typically patients with rheumatoid arthritis because of their low-demand a TWA [28]. In the past, TWA was almost exclusively used in older patients with rheumatoid arthritis because of their low-demand lifestyle, whereas patients with post-traumatic WJOA are typically young and active [22-32]. However, there is a trend in the literature towards the increasing expansion of the indication also for selected younger patients aged 31 years or younger with post-traumatic WJOA associated with good bone stock (including the Maestro™ WRS), and high claims in their activities of daily living are not always considered as contraindication [6,17,23,33,35-40]. One major complication with the use of older TWA generation types was instability or dislocation in up to 8% of cases [9,41]. It has been noted by Sterling Bunnel ([1957]: "A painless stable wrist is the key to hand function" [3]. This problem seems to be solved with the use of the new third generation types. The rates of instability or dislocation for the ReMotion™ Total Wrist is reported to be 0.7% only (1 of 144 cases), and for the Maestro™ WRS 0.6% only (1 of 157 cases) [9,22,31-34].

The non-locking Maestro™ Total Wrist, developed in 2002 by Strickland / Palmer / Graham, available in Europe since 2005 and first published with encouraging results by Dellacqua in 2009, is one of the third generation TWA that is currently in use worldwide [10,17,22,24,25,31-34,37,39,40,42-49]. The design of the implant has 3 essential differences as compared to the other third generation types (ReMotion™, Universal 2): first: the design resembles a Total Hip Arthroplasty (THA) in which a metal head articulates in a PE cup; second: the titanium alloy radial stem is to be inserted press-fit into the radial diaphysis; and third: the concave to distal shaped carpal plates are available without or with various scaphoid augments that provides a sufficient support of the carpal plate with it titanium alloy capitate peg onto the base of the trapez/trapezoid bones even in cases when the scaphoid bone is removed that is associated with the advantage that bone grafting between the surrounding carpal bones is exclusively not required (Figure 1A-C, Figure 3). The Maestro™ WRS is the further development of the implant, it has 2 changes as compared to the Maestro™ Total Wrist: first, the carpal plate is inserted with fixed or variable locking screws that should reduce the shear forces at the implant-bone interface, and in order to avoid backing out of a non-locking screw (if it would be loosened) into the articulating components’ space potentially leading to PE wear or fracture (Figure 1B); and second: the intercalated carpal heads are set up externally onto the cone of the previously inserted carpal plate whereas the carpal heads of the Maestro™ Total Wrist are screwed onto the carpal plate over the capitate peg before its insertion. This new feature allows an exchange of the carpal heads such as required in case of instability without the need for revision of the entire carpal plate (Figure 1C). Noted that the FDA in 2009 reported 1 case of backing out of a non-locking screw of the first Maestro™ Total Wrist type (inserted in this case in 2005) that required a conversion to a TWF [50].

Dellacqua noted with the first scientific paper on the Maestro™ Total Wrist in 2009 that the titanium alloy radial stem should be inserted press-fit without the need of cementation [31]. It is still widely known from literature at the beginning of the 90th of the previous century that cementation of a titanium alloy stem in THA does not confer any benefits in survivorship over its insertion in a non-cemented manner (Figure 12) [51]. Moreover, the high elasticity of titanium and the excessive stresses in the mantle of the cement could lead to micromotion and debonding of the stem, and these micromovements at the cement-stem interface may be responsible for cement mantle breakage and generation of titanium debris inducing necrosis and osteolysis [51,52]. The failure risk could be higher for smaller stems due to their greater elasticity, and in men who are physically active, while titanium stems with a larger diameter may be more successful [53]. Furthermore, Thomas et al. noted that in THA the combination of a polished titanium stem (such as with the radial stems of the Maestro™ Total Wrist/WRS) and cement appears to facilitate crevice corrosion [54]. Additionally, cementation of implants can be associated with another disadvantages such as prolonged operation time, fat embolism, and venous thromboembolic disease [55-58]. In the literature, there are no evident scientific data with the use of all third generation TWA types (including the Maestro™ Total Wrist/WRS) that support the necessity of cementation at time of its primary insertion (Figure 1A-C, Figure 5A-B, Figure 11A-E), cementation is only recommended primarily in patients with poor bone stock (rheumatoid arthritis, advanced stage of osteoporosis) (Figure 2A-B, Figure 7) or in cases of revision TWA, and cementation does not decrease the risk of intra- or postoperative periprosthetic fracture [2-13,16,17,22,49,59-63]. Noted that for metacarpal/halangeal implants it has been observed a higher risk of intraoperative periprosthetic fractures when using cementation than implants which are inserted in a non-cemented manner [64]. Moreover, the most important disadvantage of both Maestro™ types is that the PE insert is fixed to the radial body. Hence,
when PE wear or fracture occurs then a revision of the entire radial component becomes necessary. If the radial component was inserted with cementation therefore bony windowing of the distal radius shaft along the entire radial component becomes inevitable that could be associated with further complications. Both Maestro™ do not alter the Distal RadioUlnar Joint (DRUJ), hence, it can be combined with an Ulnar Head Replacement (UHR) (Figure 2B, Figure 4) [43-45,47].

An essential advantage of both Maestro™ types is the high degree of modularity that provides the combination of all components among themselves (Figure 3). Intraoperatively, the alignment and stability of implant is to be well documented by fluoroscopy especially under dynamic conditions as well (Figure 4, Figure 5A-B). An essential prerequisite in order to avoid complications such as dislocation or break out of the radial component and/or carpal plate is that both components should be exactly aligned to the axis of the radial shaft and to the axis of the capitate-3rd metacarpal bones which is the central pillar for load transfer through the wrist (Figure 6A-B), and there is a consensus in the literature for all third generation TWA types that the carpal plates should be fixed with a first screw radialwards into the 2nd metacarpal bone, and a second screw ulnarwards into the hamate bone (Figure 6C). The main complication with the use of the BIAX was break out of the metacarpal peg in dorsal direction subsequently leading to instability of its articulation and followed by PE wear and/or fracture [65]. Technical errors at time of insertion of the BIAX led to a further exacerbation of this complication rate, Takwale et al. found that for each degree a carpal component was failed placed in dorsal direction increased the risk of break out of the tip of the metacarpal peg in dorsal direction in up to 17% respectively [66]. Hence, the success for an acceptable survivorship of a TWA also depends on every surgeon’s experience. Ocampos et al. reported on their retrospective results with the use of the ReMotion™ Total Wrist (N=14), all performed by unexperienced surgeons, none of the implants were correctly aligned, and mostly observed with averaged 10,1° (range: 2°-21°) for failed positioning of the capitale peg in dorsal direction [67]. Noted that with both Maestro™ types combinations with a thumb carpometacarpal total joint replacement or a Total Elbow Arthroplasty are possible [43,47,68]. Contraindications for a TWA including both Maestro™ types are persistent osteomyelitis, irreparable nerve palsies, insufficiency of wrist stabilizing extrinsic ligaments and/or longstanding posttraumatic/postoperative wrist contracture with or without malalignment (Figure 8A-B), concomitant suspicious intraosseous lytic lesions without its diagnostic clarification, wrist hyperlaxity, untreated extrinsic (i.e. wrist-bridging) tendon disruptions, unstable soft tissue with or without infection, concomitant soft tissue tumors without
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the patient, and both for younger and elderly patients the denervation of the wrist can be successful to decrease pain over a time before TWA (Figure 10A-B) [37, 69].

Recently, in a single-center study with the use of the Maestro™ Total Wrist (published in 2015), the cumulative implant survival after 8 years (N=68) is reported to be 95%, and at the 5-year follow-up radiographic loosening was present in 2% of all cases only that is slightly superior over the other third generation types as published in 2014 [9,33]. Furthermore, the Maestro™ Total Wrist achieves the most favorable functional outcome as compared to other third-generation types, and it may be justified in preserving resection-related carpal height due to its 3 various carpal heads in combination with its design of ellipsoid surface articulation especially in order to avoid radial impingement that is mainly observed with the use of the ReMotion™ Total Wrist with its in contrast to both Maestro™ types straight design of carpal plate (Figure 1A-C, Figure 3, Figure 11A-E) [32,46,70]. However, none of all third generation TWAs are able to restore completely motion in the anatomic axes of a normal wrist including the "dart-throwing" motion arc from extension-radial deviation to flexion-ulnar deviation [71-73]. This motion arc that is performed in an oblique axis at the midcarpal joint is particularly prevalent in most occupational, recreational and avocational activities such as hammering, clubbing or fly fishing; whereas the so-called “reversed dart-throwing” motion arc from extension-ulnar deviation to flexion-radial deviation takes place almost exclusively at the radiocarpal joint [15,74]. However, a recent human cadaveric study revealed that the “dart-throwing” motion arc can also takes place at the radiocarpal joint as a compensatory mechanism when a pancarpal fusion was done [75]. In order to obtain the “dart-throwing” motion arc, a newer approach in wrist arthroplasty is midcarpal hemiarthroplasty (i.e. preserving midcarpal motion by obtaining the proximal pole of capitate bone) with the use of a radial implant alone [76,77]. However, whether this newer approach is superior to a TWA cannot be assessed currently. Additionally, another approach to obtain normal wrist circumduction with a new TWA that has some similarities with Maestro™ Total Wrist is recently in proof with a human cadaveric study [78].

Figure 5. (A) Dynamic assessment of the Maestro™ WRS intraoperatively with terminal ranges of radial and ulnar deviation (fluoroscopy) showing correct aligned implant. Note that there are no instabilities nor impingements (white arrows). (B) Dynamic assessment of the Maestro™ WRS intraoperatively with terminal ranges of flexion and extension (fluoroscopy) showing correct aligned implant in the absence of instability. Note that in this case in the presence of good bone stock the titanium alloy radial stem was inserted press-fit without the need of cementation (white arrows).

Figure 6. (A) Intraoperative clinical photograph and fluoroscopy (Maestro™ WRS) in both planes demonstrating correct central positioning of the K-wire into the radial shaft before it’s reaming. Note that for correct positioning of the K-wire in the “free hand technique” (white arrow) the axes of radial shaft are equally located in the dorsal-radial quadrant of the articular surface (green circle). (B) Intraoperative fluoroscopy (Maestro™ WRS) in both planes demonstrating reaming of the capitate bone after correct central positioning of the K-wire along the axis of the capitate-3rd metacarpal bones. (C) Intraoperative fluoroscopy (Maestro™ WRS) demonstrating placement of the 2 K-wires for fixation of the carpal plate with the first crew into 2nd metacarpal bone and the second screw into the hamate bone.

Figure 7. Same patient as in Figure 2A. The extensor retinaculum was primarily reconstructed intraoperatively, but it has disrupted 2 years after surgery again that lead to marked bowstring of the extrinsic extensor tendons (yellow arrow), and resulted in a loss of active extension at the wrist of 10° as compared with the 1-year follow up [25, 48].
Both Maestro™ types are approved for carpal hemiarthroplasty creating a “metal-on-cartilage/bone articulation” by the FDA. However, first results with a small number of patients are not encouraging. Gaspar et al. [34] reported in 2016 on 6 patients who were evaluated in mean follow-up of 35 months, and there were 1 failure, 1 instability, 3 contractures, 2 soft tissue imbalances, 2 erosions into the opposite bone, and 2 cases with DRUJ diseases. Huih et al. reported in 2017 on 11 patients who were evaluated at an average follow-up of 4 years, there was a high failure in 45% of cases that required in 2 patients a conversion to a TWA and in 3 patients a conversion in TWF, and they concluded that is probably based on a mismatch between the lunare facet of distal radius articulating with the metal head of implant [79]. Both Maestro™ types are not approved for radial hemiarthroplasty creating a “PE-on-cartilage/bone articulation” by the FDA, and the reported first results were disappointing, and it has been noted by Cooney in 2013 that care must be taken when using this implant in such an “off-label” manner [80]. The background of this procedure is similar to midcarpal hemiarthroplasty by obtaining the proximal pole of the capitate bone that allows midcarpal motion. Culp et al. reported in 2012 on 10 patients who were evaluated at mean follow-up of 19 months, and there was an unacceptable high complication rate mostly based on aseptic loosening and/or tenosynovitis (i.e. polyethylene disease) [81].

What are the causes for failed TWAs? It is still widely accepted that for older TWA types utilizing a metal-on-metal articulation that metal wear subsequently resulting in metallosis to the surrounding soft tissue, observed in nearly 100% of cases with the APH prosthesis, was the main cause for its failure [82]. Since introducing the newer TWA types utilizing a metal-on-PE articulation, metal wear appears not to be responsible for its failure. It is reported with the Universal 2 in up to 16.7%, only, and common initially presented as carpal tunnel syndrome, pisiform impingement, or pseudotumor [83-86]. With the use of the Maestro™ Total Wrist, only 1 publication could be found in which metallosis led to capsular disturbance with subsequent attenuation of the DRUJ leading to dislocation and destruction of the extensor mechanism, however, there was an impingement between the TWA and an UHR with a metal head [87]. However, the FDA reported in 2010 another case of metallosis with the Maestro™ Total Wrist due to fretting of a loosened non-locking titanium screw and the carpal plate, but a revision was required due to an infection and not due to implant issues [88]. In the literature there is a consensus with the use of the newer TWA types utilizing a metal-on-PE articulation that its failure is primarily caused by mechanical imbalance especially by the natural course of rheumatoid arthritis (ulnar carpal translocation, progressive deterioration of bone stock, muscular imbalance, ligamentous insufficiency, misalignment in the 3rd metacarpal-capitate bones axis due to angulation of the 3rd metacarpal bone in ulnar direction at the 3rd carpometacarpal joint potentially leading to unphysiological load transfer through the wrist that is observed in longstanding post-traumatic conditions as well), and secondarily followed by PE or metal wear, and noted as well that was found for the ReMotion™ Total Wrist that there are no statistically significant differences in the amount of metal or PE wear particles (evaluated in histological specimens) in patients with or without loosening of their implants [65,66,70,89,90]. The natural course of rheumatoid arthritis progressing ulnar carpal translocation especially when an ulnar head resection was performed is impressively described for dislocation of a TWA in ulnar direction, and for lunate migration onto the distal ulnar stump in patients who are not received a TWA [82,91,92]. Furthermore, the natural course of rheumatoid arthritis can also be associated with translocation of the wrist in volar direction that must be considered as contraindication for TWA as well when it leads to wrist contracture accompanied with a longstanding disruption of the volar extrinsic radioscaphocapitate ligament which is the most important stabilizer of the wrist (Figure 8A-B). However, younger patients receiving a TWA with a high mobility and functional needs may be more risky for occurrence of wear [86]. When a TWA should become necessary, correction of post-traumatic deformities at the distal radius is required in every instance if there are no post-traumatic contractures at the wrist, and pre-existent or post-traumatic WJOA is not present as well (Figure 9A-G). Furthermore, when elderly patients with advanced stage of post-traumatic WJOA are not able receiving TWA or TWF for private reasons, then the denervation of the wrist can be “last option” to improve the patient’s disability (Figure 10A-B).

What role do PeriProsthetic Osteolyses (PPO) around a TWA play in assessment of its clinical manifestation of loosening? Boeckstyns and Herzberg reported in 2014 on 44 patients received a TWA with the ReMotion™ Total Wrist and evaluated at a mean follow-up of 3.7 years, significant periprosthetic radiolucrency (more than 2 mm in width) were found juxta-articularly around the prosthetic components in 36.4% at the radial component and in 15.9% at the carpal component, whereas clinical manifestation of loosening (defined as progressive angulation or subsidence) was present in 14% of all cases only, and radiolucrency seemed to be stabilize within 3 years after surgery [93]. These observations were confirmed with the use of the Motec type in which PPO juxta-articularly around the radial component seemed to be stabilize within 8 years after surgery without any clinical signs of loosening [38]. This discrepancy between the amounts of occurrence of PPO and its real clinical manifestation of loosening is also observed in patients receiving an UHR in which PPO around the collar of implant are observed in nearly all cases and followed by its stabilization within 3 years after surgery, and that phenomenon is discussed as a result of “stress-shielding” [92]. Hence, PPOs in the absence of any clinical signs of loosening do not require surgical intervention (Figure 11A-E). That phenomenon is probably not new. Julius Wolff, a german orthopaedic surgeon (†1902), first described in 1862 that cortical bones primarily became atrophic in lesser loaded regions whereas it became hyperthrophic in higher loaded regions, and resulted secondarily in a steady-state over time (“Wolff’s law”) [95]. That phenomenon is especially observed in THA when utilizing a non-cemented titanium femoral stem which is press-fit inserted into the femoral diaphysis and followed by its sufficient osseointegration distal to the radiolucrency (Figure 12). Hence, PPOs juxta-articularly around non-cemented prosthetic components of TWAs (i.e. metaphyseal PPOs) can be probably discussed as well as result of sufficient osseointegration of its titanium alloy stems in the diaphyses distal to the metaphyseal PPOs as well.

For a failed third generation TWA there are 2 options for surgical treatment: revision TWA with or without bone grafting or cementation, and conversion to a TWF [29,60,96-100]. However, patients undergoing treatment with TWA must be prepared that it might end with TWF; therefore limited bone resection is a common feature of all contemporary wrist. Both Maestro™ implants have demonstrated uncomplicated conversion to TWF, and a recent study revealed that the outcome of TWF after a failed TWA is comparable to those after primary TWF [22,32-34,101].

**Conclusion**

TWA using all third generation types is an established motion-preserving alternative to P/TWF for treatment of WJOA. Herzberg and
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Figure 8. (A) A 73-year-old active female presented with advanced stage of rheumatoid arthritis in her right wrist. There is a marked carpal volar translocation (lines) that led to wrist contracture accompanied with longstanding disruption of the radioscaphocapitate ligament (red star). Independently that the patient is well pain-adopted with her rheumatoid-related drugs, the amount of fixed translocation of the entire carpus and the additional loss of ligamentous stabilization of the wrist are to be considered generally as contraindication for TWA. (B) Same patient with her left wrist equally to her right wrist.

Figure 9. (A) A 72-year-old active male sustained a left highly comminuted distal forearm fracture after a fall in his garden, note that there was no pre-existent WJOA. (B) Same patient, primarily surgical reduction by external fixation and pinning of the dorsally displaced extra-articular distal radius fracture, and locked plating of the distal ulna (the multiple bony fragments distal to the DRUJ were circularly sutured at the plate). The external fixation was removed 3 weeks after surgery, and the wrist was immobilized with a cast for another 3 weeks. (C) Same patient, 7 weeks after the primary injury the patient sustained a second fall-related injury associated with a left subtrochanteric femoral fracture and a re-fracture of the left distal radius with pronounced displacement in volar direction. (D) The left subtrochanteric femoral fracture was surgically reducted by a load-stable Friedl-nail. (E) Same patient, the left distal radius re-fracture was surgically treated by a correction procedure utilizing a locking distal volar radius titanium plate and a corticocancellous bone graft from the left iliac crest. Intraoperative dynamic fluoroscopy demonstrating a stable wrist at terminal ranges of passive extension and flexion. (F) Same patient 3 months later, radiographs demonstrating uneventful bony healing of the left distal radius correction procedure. (G) Same patient 6 months later, clinical photographs showing complete restoration of the active extension-flexion and supination-pronation motion arcs in comparison to the right unaffected wrist.

Adams noted in 2012: "The future appears positive for greater adoption of total wrist arthroplasty for patients who are willing to accept somewhat greater risks for the benefit of improved function" [5]. Both Maestro™ types are slightly superior in terms of modularity, survivorship at a 5 to 8-years follow-up, and functional outcome in comparison to the other types. Polished titanium alloy femoral stems (such as in both Maestro™ wrist implants) have proven to be useful and reliable in THA. In the literature, there are no evident scientific data which support the cemented use of all third generation TWAs (including both Maestro™ types!), and cementation is only recommended in the presence of poor bone stock or for revision TWA that is absolutely comparable with THA (Figure 13). Even Dr. Strickland, one of the developers of the Maestro™ Total Wrist in 2002 (!), and honoured as "Pioneer of Hand Surgery" in 2010 by the International Federation of Societies for Surgery of the Hand (IFSSH) at its 11th International Congress in Seoul (South Korea), has been noted in 2007 in original as follows: "With improved implant designs and osseointegrated stems, surgeons can achieve bone ingrowth and stability that couldn’t be achieved in the past, minimizing the need for bone cement ... In unique circumstances, however, surgeons might improve the implants’ ability by filling the medullary canal with bone cement before inserting the implant stems ... Those indications may be very poor bone stock, severe osteoporosis, previous erosive processes, or other circumstances where the surgeon feels that the stability and long-term performance of the components … may be improved by the use of supplementary bone cement ..." [102,103]. The disadvantage of both Maestro™ types is that the PE insert is fixed to the radial body, hence,
Figure 10. (A) A 72-year-old active female presented with advanced stage of right post-traumatic WJOA that allows both a TWA and a TWF. Unfortunately, she has to take care of her ill and bedridden husband after a stroke at home, and so, she was unable to endure the prolonged time for rehabilitation after TWA or TWF. Hence, the denervation of the wrist was indicated by us at last option to decrease pain in her unfortunate situation. (B) Same patient intra-operatively, the posterior and anterior interosseus nerves and the retrograde branch of the superficial radial were transected and ligated, and the neurovascular structures surrounding the radial artery were dissected carefully as well. Two months after surgery, the patient was happy that she was able to perform her activities of daily living without pain at home again.

Figure 11. (A) A 59-year-old female presented with advanced stage of WJOA accompanied with pronounced radial shortening as result of a longstanding distal radius physeal arrest and treated by a non-cemented TWA (ReMotion™ Total Wrist) primarily combined with a non-cemented UHR (metal head), formerly published by the author in 2014 at a 1-year follow-up [69]. (B) Same patient 1 year after surgery, there are no signs of PPO. Note the straight design of carpal plate of that TWA type (also at the Universal 2). (C) Same patient 2 years after surgery, first signs of PPO around the collar of UHR (yellow arrow). (D) Same patient 3 years after surgery, discrete progression of PPO around the collar of UHR (yellow arrow), and first signs of PPO at the radial component of TWA (white arrow). (E) Same patient 4 years after surgery, stabilization of PPO around the collar of UHR (yellow arrow), and progression of PPO at the radial component of TWA (white arrow) in the absence of any clinical signs of loosening, hence, revision surgery is not required at this time.

Figure 12. Long-term follow-up of THA at both sides in a 77-year-old active female who is very satisfied with her outcome. Non-cemented femoral titanium stems were inserted press-fit to the diaphyseal cortical bones (Zweymüller stems). There is a steady-state (“Wolff’s law”) between unchanged insignificant radiolucencies at the lesser loaded trochanteric regions (yellow arrows) whereas the titanium stems distal to the radiolucencies at the higher loaded regions are well osseointegrated with the cortical bones of diaphyses (white ovals).
when PE wear or fracture occurs then a revision of the entire radial component becomes necessary. When the long radial stem of both Maestro™ types is primarily inserted into the radial diaphysis with cementation, then (in case of failure) an extended bony windowing along the entire radial component is needed for its removal that can be followed by another complications in the further course. Hence, the decision by the company to withdraw both Maestro™ types from the market place because it was inserted only in a non-cemented manner in the past, is doubtful. Interestingly, Zimmer Biomet distributes 3 femoral stems (Mayo, Taperloc, M/L Taper) for THA which have similarities in surface structure (polished distally, titanium alloy) with the stem of radial component of both Maestro™ types, and all of them are approved for THAs in a non-cemented manner [104-106]. Noted that primary cementation of a TWA may also be necessary in single cases when using the ReMotion™ or Universal 2 types as a motion-preserving "off-label custom made implant" for forearm reconstruction after tumor-related large resection of the distal radius shaft [109, 110]. Finally, all surgeons who are performed TWAs utilizing both Maestro™ implants with encouraging results in the past and willing to perform it in future, now must communicate their operated and very satisfied patients as well as their new patients present with WJOA that the decision by the company Zimmer Biomet is incomprehensible and contradicts any scientific knowledge recently. Moreover, the doubtful decision by the company (i.e. no new application for CE certification which is needed for its use by the surgeons in Germany) does not allow the patients anymore to obtain an exchange TWA procedure with this implant when their previously inserted Maestro™ implants would be failed in the further course. Due to the required relatively deep bony resection into the distal radius metaphysis for placement of the radial body of both Maestro™ implants it seems to be not possible to perform an exchange procedure utilizing the other resurfacing third generation types (ReMotion™, Universal 2).

Conflict of interest

The author confirms that this article content has no conflict of interest.

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