General

Adaptive proximal scaphoid implant (APSI): a systematic review of the literature

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Scapholunate advanced collapse collapse (SLAC) is a challenging topic for hand surgeons. The adaptive proximal scaphoid implant (APSI) (Bioprofile-Tornier) is a pyrocarbon ovoid shaped interpositional implant, that allows adaptive mobility during motion. The aim of this systematic review is to analyze the clinical and radiological outcomes of APSI implants and possible complications. We performed a literature search combining the following key-words: "APSI", "Scaphoid's proximal pole", "implant", "scaphoid avascular necrosis", "SLAC", "SNAC", "pyrocarbon", "prosthesis", and "spacer" with no limitations for year of publication. We selected seven studies considered relevant to our systematic review. All studies described an improvement in the grip strength and the flexion extension arc compared to pre-operative values. The percentage of patients who reported progression of osteoarthritis (OA) with APSI was 17.3%, and implant’s mobilization has a rate 5.1% (8/156). In conclusion the APSI implant is a reliable alternative for the treatment of SNAC wrist and SLAC wrist.

INTRODUCTION

Scaphoid proximal pole nonunion treatment is a challenging topic for hand surgeons. The natural evolution will lead to osteoarthritis (OA) and scaphoid nonunion advanced collapse (SNAC).¹ Similarly, scapholunate instability leads to carpal malalignment and subsequently to osteoarthritic changes of the radio-carpal joint (scapholunate advanced collapse SLAC).² When it is not anymore possible to reconstruct the necrotic pole or to correct the carpal malalignment, salvage surgical procedures are indicated; procedures such as proximal row carpectomy (PRC), four corner fusion (4CF), or total wrist fusion (TWF) significantly decrease pain but also reduce the ROM.³

The adaptive proximal scaphoid implant (APSI) is a hydrocarbon ovoid-shaped interpositional implant that allows adaptive mobility during motion.⁴ The Pyrocarbon has good compatibility with joint cartilage and bone, a modulus of elasticity similar to bone minimizing stress shielding effects and resorption.⁵ The implant is designed with two radii of curvature: in the frontal plane, the smaller radius of curvature corresponds to the scaphoid fossa, and the larger radius of curvature is directed anteroposteriorly to the transverse plane. These two axes of the implant make it adaptable to the kinematics of the wrist.⁷ The APSI implant is an attractive solution because it is minimally invasive and does not “burn bridges” for salvage procedures such as PRC or 4CF. The implant can avoid a proximal row collapse, maintain carpal kinematics, and it is believed to prevent the progression of osteoarthritis.⁴ A systematic review was conducted to analyze the clinical and radiological outcomes of APSI implants and possible complications for evaluating if it is a safe and reliable alternative for treating SNAC and SLAC wrist.

METHODS

SEARCH STRATEGY AND LITERATURE SEARCH

A systematic review of the literature was conducted according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines.⁸ The two investigators (FS, GM) independently performed the literature search. The literature search was conducted in the following databases: Medline (PubMed), Web of Science, and Scopus were accessed on the 30th of October 2020. The following keywords were used in combination: APSI, scaphoid proximal pole, implant, scaphoid avascular necrosis, SLAC, SNAC, pyrocarbon, prosthesis, and spacer with no limitations for the year of publication. Two authors independently assessed the abstract of each publication. Article full-text was accessed for all the relevant abstracts. If the full text was not available, this warranted the exclusion from the study. The bibliography of each full-text article was also retrieved to identify additional studies.
ELIGIBILITY CRITERIA

All the studies reported data of patients undergoing the APSI procedure for SNAC and SLAC wrist. According to the authors’ language capabilities, English, Spanish, Italian, and French articles were considered. Either prospective or retrospective clinical studies were considered. Only studies published in a peer-reviewed fashion were eligible.

METHODOLOGICAL QUALITY ASSESSMENT

The two investigators (FS, GM) independently evaluated each study according to the Coleman Methodological Score (CMS).9–11 The CMS score is highly reliable, and it is widely used to assess the methodological quality of systematic reviews and meta-analyses. With this score, we analyze several characteristics of the included papers: study size, follow-up duration, surgical approach, type of study, description of surgical technique, rehabilitation, and complications. Further outcome criteria assessment, the procedure of assessing outcomes, and the subject selection process were also evaluated. CMS scores rank from 0 (poor quality) to 100 (excellent quality), with values > 60 considered satisfactory.

OUTCOMES OF INTEREST

Data extraction was performed by two independent authors (FS, GM). The following demographic data were extracted: number of patients, sex, gender, mean age, follow-up duration. The following outcomes of interest were extracted: Disabilities of the Arm, Shoulder, and Hand (DASH) questionnaire, a 10-cm Visual Analogical Scale (VAS), the Mayo Wrist Score (MWS), the Patient-Rated Wrist Evaluation (PRWEB), grip strength and Range of Motion (ROM) (flexion-extension arch), return to work, carpal misalignment, osteoarthritis progression, and complications.

RESULTS

SEARCH RESULTS

The literature search resulted in 164 articles. Of these, 144 were excluded based on the title because not inherent to our review. A further 13 articles were excluded after the reading of the abstract. Finally, seven studies were considered for the present study (Figure 1).

METHODOLOGICAL QUALITY ASSESSMENT

The CMS evidenced some limitations and points of strength of the present study. Study size and follow-up duration were poor. The surgical approach, surgical technique, and rehabilitation were well described. Complications were thoroughly discussed in most articles. Outcome measures and related timing of assessment were often defined, reporting moderate reliability. The procedure assessing outcomes and the subject selection processes were often biased and not satisfactorily described. The mean CMS was 60 (52 to 71), indicating a satisfying quality. The CMS is shown in Table 1.

PATIENT DEMOGRAPHIC

Data from 156 procedures were retrieved. 88% (126/156) were male. The average age of the patients was 45.7 years (24 to 71). The mean follow-up was 78 months (11 to 276). Demographic data are shown in Table 2.

SURGICAL TECHNIQUE

All procedures were performed using a dorsal approach except two studies: One used a volar approach in 4/41 cases and lateral in 2/41 cases, the second performed arthroscopic scaphoidectomy. Associated stioidectomy was performed in four studies for a total of 87/118 (75%) patients; two studies did not perform stioidectomy and one performed the stioidectomy, but the percentage was not reported. A dorsal capsulodesis was performed in two studies: eight of 39 patients (20%) and in all patients 36/36.

REHABILITATION

All studies had their rehabilitation program with a variable lapse of immobilization time before starting mobilization. Five studies recommended three weeks wearing full time a spica splint before beginning rehabilitation of the wrist. Aribert suggested a shorter immobilization of 2 weeks, while Gras et al. suggested an immediate ROM at home.

OUTCOMES OF INTEREST

SUBJECTIVE AND OBJECTIVE OUTCOMES

All subjective and objective outcomes are reported in Table 3. Four studies described postoperative pain using the VAS. Daruwalla reported a median preoperative VAS of 81, improving to 19 at eighteen months. The other three studies reported only the postoperative value with an average of 15.6 (range 7-22.3).

The Disability of Arm, Shoulder, and Hand score (DASH) or QDASH was evaluated in five studies. The me-
Table 1

| Part A: only one score to be given for each of the 6 sections |
|---------------------------------------------------------------|
| **1. Study size: number of patients**                         |
| <15                                                           | x   | x   |   |
| 15-24                                                         | x   |   |   |
| 25-40                                                         | x   |   |   |
| >40                                                           | x   |   |   |
| **2. Mean follow-up (months)**                                |
| <35                                                           |   |   |   |
| 36-71                                                         | x   |   |   |
| 72-107                                                        | x   | x   |   |
| >108                                                          | x   | x   | x   |
| **3. Number of different procedures included in each reported outcome** |
| Not stated                                                    |   |   |   |
| Several techniques but clearly stated                        |   |   |   |
| >1 technique but >90% receiving one technique                | x   |   |   |
| One technique                                                | x   | x   | x   |
| **4. Study type**                                            |
| Case report                                                  |   |   |   |
| Case series                                                  | x   | x   | x   |
| Retrospective comparative study                               | x   | x   | x   |
| Prospective cohort study                                      | x   | x   | x   |
| Randomized Control Trials                                     | x   | x   | x   |
| **4. Description of technique**                              |
| Inadequate/not clear                                         |   |   |   |
| Fair (technique only stated)                                 |   |   |   |
| Detailed (description of materials used)                     | x   | x   |   |
| Precise and details (picture/diagrams)                       | x   | x   | x   |
| **5. Postoperative management/rehabilitation**                |
| Not formalized                                               |   |   |   |
| Yes but unclear                                              | x   | x   |   |
| Yes and clear                                                | x   | x   | x   |
| **6. Complication discussed**                                |
| Unclear/not mentioned                                        |   |   |   |
| Mentioned but unclear                                        | x   | x   | x   |
| Fully discussed                                              | x   | x   | x   |
| **Part B: Scores may be given for each option in each of the 3 sections if applicable** |
| **1. Outcome criteria**                                      |
| ROM                                                          | x   | x   | x   |
| Further procedure                                            | x   | x   | x   |
| VAS                                                          | x   | x   | x   |

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dian postoperative DASH was 19.6 (range 7.6–26). The Mayo wrist score was evaluated in four studies\(^{2,12,13,16}\) with a median value of 74 (range 67.5–80).

All the studies measured the flexion-extension arch with an average value of 114° (range 71°–156°). The grip strength was evaluated in six studies.\(^{4,7,12–15}\)

**RETURN TO WORK**

Daruwalla et al. reported a return to work at an average of eleven weeks; Aribert et al. reported that 23/24 returned to their previous job at an average time of 2.1 months. Santos et al. reported that all patients returned to their previous job. Pequignot et al. reported that 22/25 (88%) patients returned to their previous job. Three studies did not report any information regarding return to work: Grandis et al., Poumellec et al., and Gras et al. (Table 5)

**CARPAL MISALIGNMENTS**

One study\(^{13}\) reported the presence of DISI in 46.3% of patients. One study\(^4\) did not report any changes in the SL angle. One study\(^{12}\) reported changes in the SL angle in 9/19 patients (47%). Gras et al. reported a correction of the dorsal intercalated segmental instability (DISI) in 5/8 patients (62.5%) after surgery. (Table 4)

### Table 2

| Author                  | Journal                        | Patients (n) | Mean age | Female (%) | Follow up (months) |
|-------------------------|--------------------------------|--------------|----------|------------|-------------------|
| Daruwalla et al., 2013  | Ann Acad Med Singapore        | 12           | 45 y     | 16%        | 18                |
| Aribert et al., 2019    | Hand Surg Rehabil.             | 33           | 42 y     | 6%         | 120               |
| Poumellec et al., 2019  | J Wrist Surg.                  | 19           | 42 y     | -          | 132               |
| Grandis et al., 2004    | Riv Chir Mano                  | 41           | 40 y     | 14%        | 36                |
| Pequignot et al., 2000  | Chir Main.                     | 25           | 46 y     | 4%         | 72                |
| Gras et al., 2012       | J Wrist Surg.                  | 14           | 52.7 y   | 0%         | 104               |
| Lima Santos et al., 2018| Rev Bras Ortop.                | 12           | 39 y     | 0%         | 66                |

**PROGRESSION OF OSTEOARTHRITIS**

One study\(^{13}\) reported a progression of osteoarthritis in 6/35 patients (18%). Poumellec et al. report a progression of the OA in 6/19 patients (32%). Gras et al. reported a radial styloidectomy 2-5 years after the APSI procedure in 3/14 patients (21%) for radio-scaphoid arthritis progression. Santos et al. reported OA progression in all the cases (12/12). Three studies\(^4,14–16\) did not report any OA progression. The percentage of patients who reported progression of OA was 17.3% (27/156; Table 4).

**COMPLICATIONS**

Four studies reported no complication.\(^4,14–16\) Gras et al. reported two mobilizations of the implant treated with a 4CF. Poumellec et al. reported two cases of early implant dislocations due to a lack of compliance to the postoperative immobilization required. Aribert et al. described 6/33 complications (18%). One case of early sepsis required implant removal, four instances of dislocation, and one patient complained about persistent pain caused by radio-scaphoid impingement and was treated with a 4CF. The total amount of mobilization of the implant was 5.1% (8/156; Table 4).
### Table 3

| Author                  | Return to work | DASH Pre op | DASH Post op | MWS Pre op | MWS Post op | POWER Pre op | POWER Post op | VAS Pre op | VAS Post op | Grip strength | ROM Pre op | ROM Post op |
|-------------------------|----------------|-------------|-------------|------------|-------------|--------------|--------------|------------|-------------|---------------|------------|-------------|
| Daruwalla et al.        | 11 w           | 55          | 20          | -          | -           | 81 (0-100)   | 19 (0-100)   | 30 kg      | 30 kg       | 73°           | 80°        |
| Aribert et al.          | 8 w            | -           | 19.5        | -          | 80          | 17.5         | 1.2 (0-10)   | 27 kg      | 38 kg       | 99°           | 101°       |
| Poumellec et al.        | -              | 26          | -           | 69         | -           | 25           | -            | 34 kg      | -           | 106°        |
| Grandis et al.          | -              | -           | -           | -          | -           | -            | -            | -          | 40% of improvement | -           | 100°       |
| Pequignot et al.        | -              | -           | -           | -          | -           | -            | -            | -          | 80% of controlateral | -           | 95°        |
| Gras et al.             | -              | 7.6         | -           | 79.6       | -           | 7.5 (0-10)   | 0.7 (0-10)   | 15.8 kg    | 44.1 kg     | 77°           | 136°       |
| Lima Santos et al.      | -              | 25          | -           | 67.5       | -           | -            | -            | 22.3 kg    | -           | 71°          |

### Table 4

| Authors                  | Numbers of patients | Osteoarthritis progression | Malalignment | Complications |
|--------------------------|---------------------|---------------------------|--------------|---------------|
| Daruwalla et al          | 12                  |                           | 41.6% of DISI | 4 dislocations |
| Aribert et al.           | 33                  | 6                         | 47% of DISI  | 1 impingement  |
| Poumellec et al.         | 19                  | 6                         | 8% of DISI   | 2 dislocations |
| Grandis et al.           | 41                  |                           |              |               |
| Pequignot et al.         | 25                  |                           |              |               |
| Gras et al.              | 14                  | 3                         |              |               |
| Lima Santos et al.       | 12                  | 12                        |              |               |
DISCUSSION

Some of the most widely used treatment options for wrist pain include PRC and 4CF. A systematic review in 2009\(^3\) compared these two techniques and found no differences in terms of pain, grip strength, and subjective outcome. These data suggest that PRC has fewer potential complications and better ROM than 4CF but a higher risk of progression of OA. Some researchers have tried to replace the scaphoid with interposition of autologous material,\(^17,18\) silicon,\(^19\) or metal,\(^20\) but the results were not satisfactory.

The average quality of the studies included in the present investigation is moderate, with an average CMS of 60 points; only one study\(^15\) had good CMS values. The most common surgical approach was from the dorsal wrist except for two studies that adopted an arthroscopic approach.\(^7,15\) Arthroscopy is an attractive option for its minimally invasive procedure and consequent preservation of soft tissues, including ligaments and capsules.

Stiloidectomy is widely performed\(^4,12–15\) (73% of all patients) and is advisable in the case of a stylo-carpal impingement. Two papers\(^12,13\) performed a dorsal capsulodesis with no improvement in implant stability. There were six dislocations of the implant out of eight in these studies. All studies reported a postoperative immobilization of 2-3 weeks except for one,\(^7\) which allowed an early ROM probably due to the low invasiveness of the arthroscopic procedure.

All the studies described a reduction in grip strength and the flexion-extension arch compared to the contralateral wrist but with improvement versus preoperative values. The highest value in flexion-extension arch (136° vs. an average value of 92°) and grip strength (44.1 kg vs. an average value of 54 kg) were reported by Gras et al., who performed the replacement arthroscopically and with an early rehabilitation protocol. The same study\(^7\) reported the best score in subjective outcomes such as DASH and VAS. We hypothesize that this is due to less scar tissue and a more aggressive rehabilitation protocol.

Gras et al. reported a correction of the dorsal intercalated segmental instability (DISI) in 5/8 patients (62.5%) after surgery. The role of the APSI is controversial in its impact on the DISI deformity: two studies\(^12,13\) reported an increased SL angle in 46% of patients despite the use of capsulodesis. We speculated that the arthroscopic approach preserves the secondary stabilizer versus the open technique.

The percentage of patients who reported progression of OA with APSI was 17.3%, the average increases to 26% (27/105) if we consider only studies with more than five years in the follow-up: Aribert et al., Poumellec et al. Pequignot et al., Gras et al. Santos et al. The APSI slowed the progression of OA but did not stop it. The implant mobilization has a rate of 5.1% (8/156). The surgical approach and postoperative immobilization does not correlate with this value. In one series,\(^7\) the use of the implant for a nonunion of the body of the scaphoid leads to mobilization.

All procedures were performed using a dorsal approach\(^4,12–14,16\) except two studies: One used a volar approach in 4/41 cases and lateral in 2/41 cases,\(^15\) the second performed arthroscopic scaphoidectomy.\(^7\) Associated stiloïdectomy was performed in four studies\(^4,12,15,16\) for a total of 87/118 (73%) patients; two studies did not perform stiloïdectomy,\(^7,16\) and one\(^14\) performed the stiloïdectomy, but the percentage was not reported. A dorsal capsulodesis was performed in two studies—eight of 39 patients (20%)\(^15\) and all patients 36/36.\(^12\)

All studies had their rehabilitation program with a variable lapse of immobilization time before starting mobilization. Five studies recommended three weeks wearing full time a spica splint before beginning rehabilitation of the wrist.\(^4,12,14–16\) Aribert suggested a shorter immobilization of 2 weeks, while Gras et al. suggested an immediate ROM at home.

The study has several limitations. For example, we grouped different cohorts of patients in terms of demographics, surgical indication, surgical technique, outcome measurements, and follow-up.

The main endpoint of this systematic review is that the APSI implant is a reliable alternative for treating SNAC wrist and SLAC wrist. Its indications are limited: The cartilage of the scaphoid fossa and the capitulonate joint must be intact. However, the implants require less surgical dissection compared to 4CF and PRC. This procedure is more expensive than 4CF and PRC due to the high cost of the spacer, but further surgeries are possible in case of failure.

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