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OFFICE OF THE PUBLIC AULITOR
PROCUREMENT A PYRALS

UCI 24 2008 TIME: 20194

OFFICE OF THE PUBLIC AUDITOR FILE No. OPA-P.

FILE No. OPA-PA - 07-01/

IN THE APPEAL OF

Facsimile: 671-475-8550
Attorneys for Appellee GMHA

JMI MEDICAL SYSTEMS, INC.

Appellant.

Appeal No. <u>OPA-PA-07-011</u>

GMHA'S RESPONSE TO JMI'S MOTION FOR SUMMARY JUDGMENT

Comes now, Appellee Guam Memorial Hospital Authority (hereinafter referred to as "GMHA") who opposes JMI's Motion for Summary Judgment. The Summary Judgment standard states that the Judgment shall be rendered forthwith if ... there is no genuine issue as to any material fact and that the moving party is entitled to judgment as a matter of law. The Appelle, GMHA, respectfully contends that there is a genuine issue as to material facts present.

The Motion for Summary Judgment pursuant to Rule 56 is seriously flawed. First, it is asserted that there are no issues or facts in controversy and that Summary Judgment should be issued forthwith. On the contrary, witness Jean Grape Ko, a key employee of Appellant, testified that JMI was not sure what the specifics were for bidding on the test kits. *See* Deposition Transcript of Jean Grape Ko (hereinafter "Ko Deposition") at page 19, attached hereto as Exhibit A.

As an aside, Appellee, for the record, respectfully disagrees that 2 GAR § 12104 (c)(4) is controlling procedurally in this matter as to the scheduling of its deadline to file this opposition. That provision deals with comments on the agency record and the agency's response to said comments and is not applicable to the type of motion filed by Appellant. Furthermore, the OPA rules and regulations do not address this issue. Appellee was given two (2) days notice of when its Opposition was due. We agree with the Hearing Officer's determination that the Guam Rules of Civil Procedure do not apply as administrative process is governed by 2 GAR, Division 4, Chapter 12 Rules for Procurement Appeals. However, Appellee was unsure of what time period applied to the filing of its opposition and because it was Appellant who propounded this motion for summary judgment pursuant to the GRCP Appellee assumed it had 21 days to respond considering Appellee did not receive notice from the Hearing Officer of a different time within which it was to file an opposition or response to Appellant's motion.

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Ms. Grape Ko testified that she did not know what the bid prices for the test kit prices were but she totally depended on the manufacturer's suggested prices and types of test kits. See Exhibit A. The manufacturer did not specifically research nor did he know which test kits were needed by Appellee GMHA. See Ko Deposition at pages 25-26, attached hereto as Exhibit B. Neither Ms. Grape Ko nor any of the three witnesses deposed by Appellee GMHA were aware of which test kit they were bidding on.

Second, Appellant argues that Appellee violated its own procurement regulations by purchasing a second analyzer and supplies without going through competitive sealed bidding as required by law. Put another way, Appellant argues that there is a second procurement of machines under the initial procurement. This is the very first time that such an issue has been raised. Appellant did not include this argument in its initial protest with the OPA and in subsequent filings. Appellant should not be allowed to further this argument as it has failed to give Appellee sufficient notice such that Appellee can properly prepare defenses to this argument and Appellant has waived the ability to set forth this argument at this late date. It is rather disingenuous, to bring it up at this time, since it is a new issue and this appeal is based on the issues that have raised.

Third, Appellant concedes the fact that it submitted a bid that was extremely high for each test kit as compared to the test kit prices submitted by MedPharm. Appellant's submission of a high bid was after the request had been made for the bid prices on the specific kits by GMHA. As MedPharm's bid specifications met GMHA's requirements, MedPharm was the successful bidder. Appellant exhaustively argues that its bid complied in all material aspects to the Invitation for Bid. In light of Appellant's concession that its bid was too high, whether Appellant's bid was materially compliant is clearly in dispute.

Fourth, JMI's interprets events to suit its needs. JMI says that the bidder submissions were wrongfully considered in that the additional cost information by MedPharm and also

wrongfully considered the ability of GMHA to consider the purchase of not one but two analyzers for the purchase of one. This is a new assertion. We can posit a different scenario having occurred.

Fifth, under paragraph 25 of the Solicitation, GMHA can unquestionably reject bids in parts. As there is an obvious conflict between paragraph 7 and 25, such a conflict is nonetheless legally resolvable. There is a principle of contract law which states that: "where repugnancy is found between clauses one which essentially requires something to be done to effect the general purpose of the contract it is entitled to greater consideration than the other". 17 Am. Jur. 2d Contracts § 384 citing International Union of Operating Engineers v. J.A. Jones Const. Co., 240 S. W. 2d 39 (Ky. 1951).

The Procurement Law allows the application of the above described contract law principle. See 5 GCA § 5002 ("the principles of law and equity including the Uniform Commercial Code, the law merchant shall supplement" the Guam Procurement Law.") Thus, the rejection of the "reagent & supplies" part of the bid submitted is legally permissible and should be upheld. GMHA has acted in good faith in handling the procurement in issue.

In summary, Appellant's motion should be DENIED as there are genuine issues as to material facts of which Appellant relies upon in bringing its motion for summary judgment.

Dated this 24th day of October, 2008.

The Law Offices of John S. Unpingco & Associates, LLC

By:

John S. Unpingco, Esq

which will be after the bid evaluation of the user. They will identify which specific test they will provide.

- Q. So you're basing the clarity of the specification just on the number of tests to be done per annum?
- A. Yes, and that we asked our vendor, because they are the one, the expert, to identify how many kits that will be used in 3,000 tests.
- Q. Okay. By vendor, you mean like Vitek, the manufacturer of the machine?
- A. The dealer of the Biomerieux. Biomerieux is the manufacturer of Vitek.
 - 💮 Q. Biomer- --
 - A. Biomerieux.
 - Q: How do you spell that?
- A. B-I-O-R-E-I-M-E-U-I-X. Like that. Oh! B=I=O-M-E-R-I-E-U-X.
- So is it fair to say if you're just going by the number of tests alone, that does not tell you the specific type of tests that are to be performed, just the number?
- O. Is it fair to say that when you're looking at that requirement for 3,000 tests to be done per year, it's only telling you how many tests are to be done, it doesn't tell you the exact test that are to be performed?
 - A Mes, sir, because it's stated that to supplies for

based on the specs given and the award is not fair due to the fact that the other vendor -- Can I say the name?

Q. Yes.

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- Which is Med Pharm, is nonresponsive by not offering the reagents or the kits for the 3,000 tests per annum.
- But if your offer did not specify which test you're Q. offering, are you not -- is not your bid also nonresponsive to what the hospital needs?

MR. SISON: Well I'm going to object to that as it calls for legal conclusion, speculation.

MR. UNPINGCO: Objection noted but I think as a matter of common sense, I would like to find out her opinion whether or not she felt that it was meeting the specifications.

> THE WITNESS: Should I answer?

MR. SISON: Answer, as you can.

THE WITNESS: Okay. Well in my understanding, to off a general terms of the specific kits, like the GPS and GNS, the lab tech or the supervisor understand that kind of general terms for that and they will provide whichever the vendor offered this particular award and they will provide which test they will use.

- (By Mr. Unpingco) So there's still a missing piece Q. of information?
 - Α. Yes, and in my opinion, this bid should compose on

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that from the hospital. They should notify which specific test they want.

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- Q. So their specs were ambiguous, because they -- it was not complete?
- A. I don't think it's ambiguous, because they said to provide tests or supplies, then we ask our vendor to provide the test or the kits for that 3,000 test per annum because they're our expert on that particular area, on that laboratory equipment and they can provide how many kits on the 3,000 tests per annum.
- Q. But it was not the vendor that was purchasing the kits. The problem I guess I'm struggling with here is that you satisfied the requirement for 3,000 tests, that I can see you doing, you're turning to your vendor, but the other part is of those 3,000, are those the correct 3,000 that the hospital is seeking, and what I'm asking you is that the hospital, and I think you've answered that already so forgive me if I'm repeating it, but the hospital never specified what particular test those 3,000 were to consist of.

They never said a thousand tests of this particular type. Until the August 10th letter, that's when you first received an idea of what test, which specific tests the hospital was after. Is that fair to say?

- A. Yes, sir, that's after.
- Q. After the fact, after the bid opening. Right?