

Injection Force of SoloSTAR[®] Compared with Other Disposable Insulin Pen Devices at Constant Volume Flow Rates

Thomas van der Burg, B.Sc.

Abstract

Background:

Injection force is a particularly important practical aspect of therapy for patients with diabetes, especially those who have dexterity problems. This laboratory-based study compared the injection force of the SoloSTAR[®] insulin pen (SoloSTAR; sanofi-aventis) versus other available disposable pens at injection speeds based on the delivered volume of insulin released at the needle.

Method:

Four different prefilled disposable pens were tested: SoloSTAR containing insulin glargine; FlexPen[®] and the Next Generation FlexPen[®] (NGFP) (Novo Nordisk), both containing insulin detemir; and KwikPen[®] containing insulin lispro (Eli Lilly). All pens were investigated using the maximum dispense volume for each pen type [80 units (U) for SoloSTAR; 60 U for the other pens], from the free needle tip dispensing into a beaker. Twenty pens of each type were fitted with the recommended needles and tested at two dose speeds (6 and 10 U/s); each pen was tested twice.

Results:

Mean plateau injection force and maximum injection force were consistently lower with SoloSTAR compared with FlexPen, NGFP, and KwikPen at both injection speeds tested. An injection speed of 10 U/s was associated with higher injection force compared with 6 U/s for all the pens tested ($p < .001$).

Conclusions:

SoloSTAR stands out because of its low injection force, even when compared with newer insulin pen devices such as the KwikPen and NGFP. This may enable patients, especially those with dexterity problems, to administer insulin more easily and improve management of their diabetes.

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Author Affiliation: sanofi-aventis Deutschland GmbH, Site Frankfurt Devices/Device Industrialisation, Frankfurt, Germany

Abbreviations: (N) newton, (NGFP) Next Generation FlexPen, (U) unit

Keywords: flow rate, injection force, injection speed, insulin pen

Corresponding Author: Thomas van der Burg, sanofi-aventis Deutschland GmbH, Site Frankfurt Devices/Device Industrialisation, Industriepark Hoechst, D-65926 Frankfurt am Main, Germany; email address thomas.vanderburg@sanofi-aventis.com

Introduction

The first insulin pen device was introduced in 1985. Since then, continuing innovation has led to a steady improvement in the devices available and they now account for about half of worldwide insulin use.¹

There are numerous disposable pen devices available on the market in the United States,² Europe, and Japan, such as FlexPen® (Novo Nordisk, Bagsvaerd, Denmark), more recent disposable devices such as KwikPen® (Eli Lilly, Indianapolis, IN), and the so-called Next Generation FlexPen® (NGFP) (Novo Nordisk, Bagsvaerd, Denmark).

The Lantus® SoloSTAR® disposable injection device (sanofi-aventis, Paris, France) was launched in 2007 and meets a combination of user needs that had not been previously addressed and still remain unmet by other devices on the market. These include ease of injection, differentiation of insulin type through pen body color and tactile elements, and the ability to inject up to 80 units (U) of insulin in one injection with a comparatively short dial stroke, which is particularly useful for patients with impaired manual dexterity.³ The SoloSTAR pen was developed through a process of iterative design and feedback questionnaires involving patients, healthcare professionals, the design team, and consultants in order to comprehensively assess the needs of patients who use insulin pens.³

Injection force is a particularly important practical aspect of therapy for patients with type 1 or 2 diabetes, especially for those who have dexterity problems; these patients may have limited ability to self-inject insulin.⁴⁻⁷

The aim of this study, therefore, was to compare the injection force of the SoloSTAR pen with three other commonly available disposable pens at two different injection speeds based on a delivered volume of insulin released at the needle (constant volume flow rate) within a laboratory setting. This is the first study directly evaluating the injection force of these three insulin devices on the basis of the dispensed dose per time, using realistic dispense speeds for practical use.

Methods

Four different pen injection devices were tested in this investigation: SoloSTAR insulin glargine pen (batch number 40U286), FlexPen insulin detemir pen (batch

number VH70215), NGFP insulin detemir pen (batch number VH70007), and KwikPen insulin lispro pen (batch number A477063).

Twenty pens of each type were tested at two dose speeds (6 and 10 U/s); each pen was tested twice, with all doses delivered into a beaker. Tests were carried out using the maximum dial stroke and dispensing the maximum dose volume of each pen type (80 U for the SoloSTAR; 60 U for the comparator pens). All investigations were conducted using the manufacturers' recommended needles with a consistent outer diameter of 0.25 mm based on the manufacturers' specifications: BD Micro-Fine 0.25 mm (31G) × 5 mm for SoloSTAR and KwikPen; NovoFine 0.25 mm (31G) × 6 mm for FlexPen and NGFP.

Laboratory tests were conducted using a tensile meter (Zwick GmbH & Co. KG, Ulm, Germany) and force cell [KAF-TC, Zwick GmbH & Co. KG, Ulm, Germany; nominal load 200 newtons (N)] under standard atmospheric conditions. The distance traveled by the push button to deliver the appropriate dose was determined to be different for each of the pens, necessitating a different push button speed to be chosen for the four devices. Before evaluating each pen, the appropriate needle was mounted and correct fitting ensured by dispensing a priming dose of 10 U. For each pen device, the injection force throughout the dose delivery was measured (**Figure 1**). The mean force value (mean plateau injection force) was calculated and the maximum injection force evaluated.

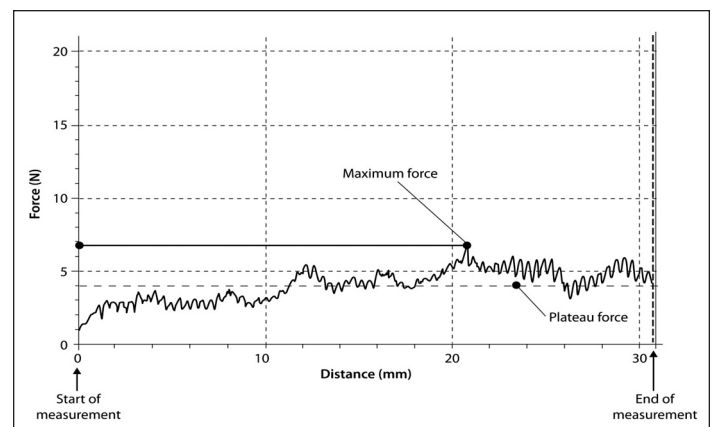


Figure 1. Example force measurement curve. Laboratory tests were carried out in order to determine the injection force of insulin pen devices at maximum insulin dose and two injection speeds (6 and 10 U/s). Injection force throughout the dose delivery was measured and mean plateau and maximum values evaluated.

Statistical analyses were carried out using Dunnett's test. A simultaneous test level of $p < .05$ was assessed with SoloSTAR as the reference group; the corresponding individual test level was $p < .012$ for each of the compared pairs. The differences in the mean maximum injection force and the mean plateau injection force between SoloSTAR and FlexPen, NGFP, and KwikPen were calculated, and the resultant confidence intervals were determined. The compared pair can be assumed to be different ($p < .012$), if the confidence interval of the respective pair is larger than 0. Due to the sufficient level of significance, no further declaration for the probability for the tested pairs was made.

Preliminary studies revealed that the injection force of insulin devices with the needle attached is mainly determined by the following factors: friction between the mechanical parts of the mechanism; friction between the bung and the glass partition of the cartridge; fluid friction of the liquid (insulin); and tissue pressure. Because tissue pressure is difficult to measure or simulate with high consistency, all tests were performed by dispensing into a beaker. Preliminary studies also showed that the fluid friction of the expelled insulin is mainly affected by changes in flow rates. Therefore, only comparisons at equal volume flow rates were pursued in this study.

Theoretical considerations were made to determine the dependence of fluid friction on the volume flow rate. Using basic fluid dynamics theory, one can demonstrate that the fluid friction of the insulin being expelled causes an accessory pressure inside the cartridge that increases the force required by the user to dispense the dose. The additional pressure can be calculated using the formula in Equation (1), which is derived from the Bernoulli equation⁸ by adding terms for the pressure reduction caused by fluid friction⁹ and cross-section changes.⁸ In Equation (1), ρ represents the density of the fluid, d_{needle} the inner diameter of the needle, l_{needle} the total length of the needle, \dot{V} the volume flow rate; α and ξ_2 are empiric coefficients mainly caused by crossover at the needle tip, ξ_1 is an empiric coefficient for the crossover of the fluid between cartridge and needle, and λ is the coefficient of friction for the needle (depending on viscosity, flow rate, roughness of the needle, and needle diameter).

$$\Delta p = \frac{8 \cdot \rho}{\pi^2} \cdot \dot{V}^2 \cdot \left[\left(\frac{\alpha}{d_{needle}^2} \right) + \frac{1}{d_{needle}^4} \left(\lambda \cdot \frac{l_{needle}}{d_{needle}} + \xi_1 + \xi_2 \right) \right] \quad (1)$$

As ρ , d_{needle} , l_{needle} , α , ξ_2 , ξ_1 , and λ are roughly constant for one needle/device combination at the used flow rate area, they can be expressed as the constant coefficient B , resulting in the simplified formula in Equation (2), where the accessory pressure inside the cartridge only depends on the volume flow rate of the insulin.

$$\Delta p = B \cdot \dot{V}^2 \quad (2)$$

This formula was used to verify the theoretical approach by calculating the increase of the injection force at 10 U/s compared with 6 U/s for SoloSTAR with BD 0.25 mm (31G) × 5 mm needles. Subject to λ , an increase in the range of 2.8–4.4 N could be expected because of the higher volume flow rate.

Results

The mean plateau injection force at the maximum doses with the pens (80 U for SoloSTAR versus 60 U for the comparator pens) was significantly higher with FlexPen, NGFP, and KwikPen compared with SoloSTAR at both injection speeds tested (Figures 2A and 3A). The difference in mean plateau injection force compared with SoloSTAR for the various pens was 95, 51, and 43% with FlexPen, NGFP, and KwikPen, respectively, at 6 U/s, and 87, 47, and 37%, respectively, at 10 U/s (Table 1). An injection speed of 10 U/s was associated with higher injection force compared with 6 U/s in all the pens ($p < .001$).

In line with the mean plateau force, the maximum injection force was also significantly higher with FlexPen, NGFP, and KwikPen compared with SoloSTAR at both injection speeds tested (Figures 2B and 3B). The difference in maximum injection force compared with SoloSTAR for the various pens was 70, 26, and 29% with FlexPen, NGFP, and KwikPen, respectively, at 6 U/s, and 65, 31, and 30%, respectively, at 10 U/s (Table 1).

FlexPen showed the highest injection forces of all tested devices. Although KwikPen and NGFP showed comparable maximum forces, the mean plateau force of KwikPen was calculated to be significantly lower than that of NGFP ($p < .012$).

Discussion

Among the four disposable insulin pen devices compared in this study, the SoloSTAR pen had the lowest injection force irrespective of the injection speed tested. The dispense force of all pens rose when dispensing the dose at higher

speed, but injection force for SoloSTAR remained significantly lower than those of the other pens. This difference was observed for both mean plateau force and maximum force.

The empirical increase of the injection force at 10 U/s compared with 6 U/s for SoloSTAR was 3.7 N for the mean plateau force, which corresponds to the theoretical expectation with a calculated increase in the range of

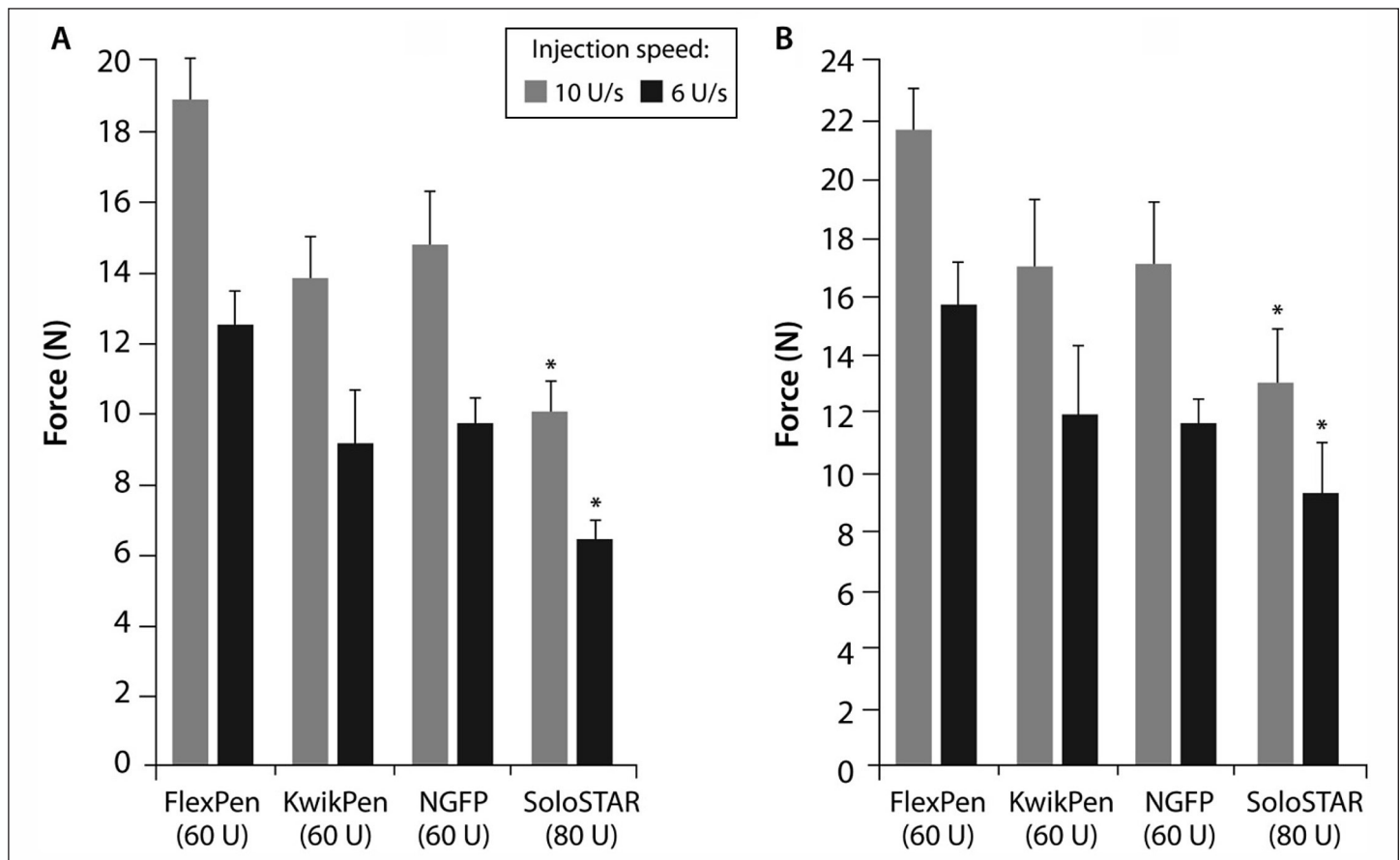


Figure 2. Comparison of mean plateau injection force (A) and maximum injection force (B) at two injection speeds for each pen and dose tested. Mean plateau and maximum injection forces of the various insulin pen devices were measured at maximum insulin dose for each pen (80 U for SoloSTAR; 60 U for the other pens and at two injection speeds (6 and 10 U/s) for each pen. Twenty pens of each type of device were tested twice for each dose and speed combination, and average values calculated. * $p < .001$ compared to comparator pens at the same injection speed.

Table 1. Injection Force with Various Disposable Insulin Pen Types, Doses, and Injection Speed Combinations

	Dose (U)	Injection speed (U/s)	Button speed (mm/s)	Mean plateau injection force \pm SD ^a (N)	Min–Max values (n)	Maximum injection force \pm SD ^a (N)	Min–Max values (n)
SoloSTAR	80	6	2.6	6.43 \pm 0.59	5.22–7.60	9.30 \pm 1.71	6.17–14.68
	80	10	4.3	10.10 \pm 0.84	8.72–12.15	13.10 \pm 1.90	10.24–17.93
FlexPen	60	6	3.3	12.51 \pm 0.96	10.72–14.30	15.79 \pm 1.41	13.07–19.57
	60	10	5.5	18.91 \pm 1.24	16.05–20.91	21.64 \pm 1.52	18.46–23.72
NGFP	60	6	3.3	9.72 \pm 0.72	8.44–11.87	11.71 \pm 0.83	10.32–13.75
	60	10	5.5	14.79 \pm 1.50	12.10–18.92	17.13 \pm 2.17	13.76–25.30
KwikPen	60	6	2.8	9.17 \pm 1.54	6.54–12.50	11.95 \pm 2.52	6.67–16.58
	60	10	4.7	13.82 \pm 1.31	11.32–16.35	16.99 \pm 2.40	13.13–22.51

^a SD = standard deviation

2.8–4.4 N. Therefore, the results of this study confirm the theoretical approach that the volume flow rate provides the main influence on the injection force for a specific pen/needle combination.

The findings from this study are in line with previously published laboratory-based studies in which the SoloSTAR pen had an improved injection force compared with the FlexPen device.³

The findings of this study disagree with the results of one study by Rissler and colleagues¹⁰ and one study by Asakura and colleagues,¹¹ which suggested that the NGFP had a lower injection force compared with SoloSTAR. To understand the relevance of these conflicting data, it is important to recall that constant volume flow rates (U/s) were used in our study, whereas the other two studies used different injection button speeds (mm/s). The latter methodology means that, even at equal injection button speeds, the insulin flow in terms of U/s is not the same between the pen systems. Owing to the differences between the mechanisms of the pens, the volume of insulin expelled per second for SoloSTAR is 27.5% larger than that for FlexPen and NGFP. Elucidating the differences in the volume flow rates, dispensing the same volume of insulin with SoloSTAR requires a 22% smaller push-button travel and injection time (not accounting for differences in the holding time). This shorter push-button travel as the result of the shorter dial stroke extension is likely to be preferable for patients with impaired dexterity^{3–7,12} as well as unimpaired patients.^{13,14}

An observational, survey-based clinical study by Carter and colleagues reported high levels of acceptance of the SoloSTAR device among patients both with and without manual or dexterity impairment.¹³ Participants found SoloSTAR easy to use and that using the pen had a positive impact on the management of their diabetes, such as increasing confidence and helping overcome their reluctance to use insulin.¹³ In a study by Haak and colleagues where 16% of patients had dexterity problems and 19% visual impairment, more patients preferred the effort required to inject a 40 U dose with SoloSTAR versus FlexPen.¹⁴ The findings of these studies may relate to the lower injection force characteristics of SoloSTAR versus the FlexPen device as demonstrated by Clarke and Spollett.³ However, both of these studies demonstrated the usability and acceptance of SoloSTAR in these populations; they did not report the impact of injection force on the outcomes. Therefore, prospective studies are needed to extend these findings in patients with and without dexterity problems, and investigate whether

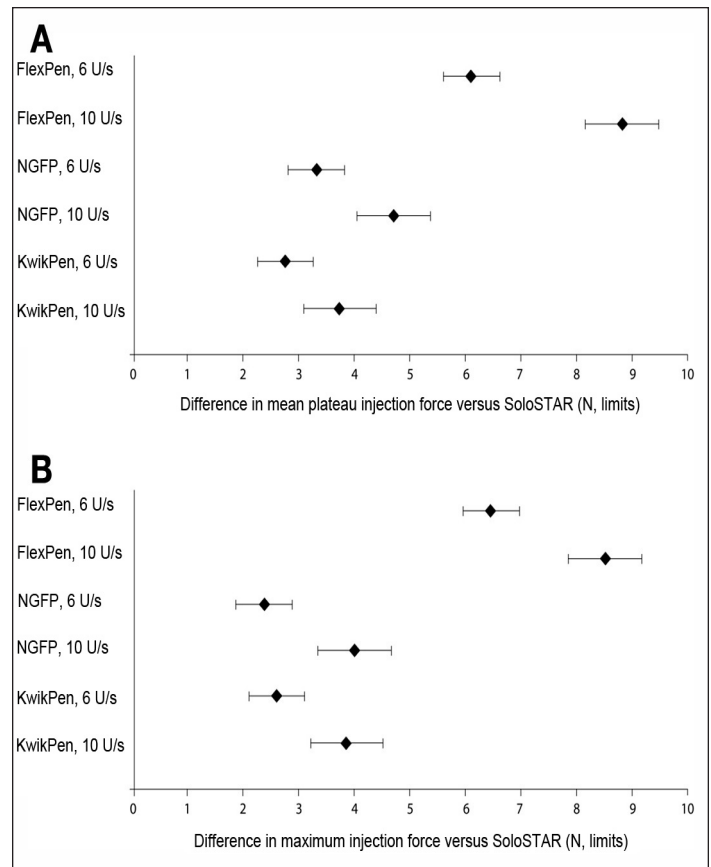


Figure 3. Interval of differences in mean plateau injection force (A) and maximum injection force (B) between SoloSTAR and the various comparators at maximum doses tested. Mean plateau and maximum injection forces of the various insulin pen devices were measured at maximum insulin dose for each pen (80 U for SoloSTAR; 60 U for the other pens) and at two injection speeds (6 and 10 U/s) for each pen. The average values for 20 pens tested twice each were calculated, and hence the difference between the average forces for SoloSTAR and each of the comparator pens as well as the limits (calculated with Dunnett's test for the difference between the respective group mean and SoloSTAR at 80 U). The differences in the mean maximum injection force and the mean plateau injection force between SoloSTAR and FlexPen, NGFP, and KwikPen were calculated with the resultant confidence intervals shown in Figures 3A and B. Only if 0 is within the confidence interval of the respective pair does the tested device show no significant difference to SoloSTAR. If 0 is not within the confidence interval, the compared pair can be assumed to be different with a probability of at least 98.8%. Due to the sufficient level of significance ($p < .012$), no further declaration for the probability for the tested pairs seems to be of value.

the low injection force of available insulin pens is an important factor in their use of insulin and ultimately diabetes management.

In order to maximize clinical relevance, we evaluated each pen together with the manufacturer's recommended needle in order to emulate real-world use. While needle outer diameters were consistent, potential variations of the inner diameter of needles within the manufacturers' specifications may have contributed to differences observed between the pens. It must also be acknowledged that

the study was performed in a simulated, laboratory environment, with doses delivered into a beaker rather than injected into tissue. Therefore, the data may not fully reflect patients' perceptions of the forces encountered in everyday use of the devices because the injection force may be affected by the different properties of the tissue being injected into and the individual characteristics of the injection. Further investigations using more than the two different dispense speeds tested in this study might also be worthwhile to verify the influence of the volume flow rate. Additionally, the impact of injection force and injection speed on patients' perceptions of the pain of injection should be evaluated because there are limited data at present. A rapid speed of injection may be preferable to patients but may be achieved at the expense of increased injection force and increased patient discomfort.

Conclusions

The results of this study confirm the theoretical approach that the injection force for a specific pen/needle combination is mainly influenced by the volume flow rate. Therefore, it can be said that comparisons of injection forces at fixed button speeds are misleading and that the methodology used in our study provides a more realistic picture of the performance characteristics of the pens tested.

The mean plateau injection force as well as the maximum injection force was significantly lower with SoloSTAR compared with FlexPen, NGFP, and KwikPen at both injection speeds tested. The lower injection force needed with SoloSTAR versus the comparator pens may have a positive impact on the management of diabetes, particularly in patients who have dexterity issues.

Even compared with newer insulin pen devices such as KwikPen and NGFP, SoloSTAR stands out due to its low injection force as well as the possibility to inject up to 80 U of insulin in one injection with a comparatively short extension of the dial stroke.

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Disclosure:

Thomas van der Burg is an employee of sanofi-aventis.

References:

1. IMS Health. IMS Midas™ June 2006, Quarterly insulin sales volume in units. 2006.
2. Selam JL. Evolution of diabetes insulin delivery devices. *J Diabetes Sci Technol*. 2010;4(3):505-13.
3. Clarke A, Spollett G. Dose accuracy and injection force dynamics of a novel disposable insulin pen. *Expert Opin Drug Deliv*. 2007;4(2):165-74.
4. Rosenbloom AL. Limitation of finger joint mobility in diabetes mellitus. *J Diabet Complications*. 1989;3(2):77-87.
5. Savaş S, Koroğlu BK, Koyuncuoğlu HR, Uzar E, Celik H, Tamer NM. The effects of the diabetes related soft tissue hand lesions and the reduced hand strength on functional disability of hand in type 2 diabetic patients. *Diabetes Res Clin Pract*. 2007;77(1):77-83.
6. Schady W, Abuaisha B, Boulton AJ. Observations on severe ulnar neuropathy in diabetes. *J Diabetes Complications*. 1998;12(3):128-32.
7. Ziegler D, Gries FA, Spüler M, Lessmann F. The epidemiology of diabetic neuropathy. DiaCAN Multicenter Study Group. *Diabet Med*. 1993;10(Suppl 2):82S-6S.
8. Kast W. Druckverlust in Leitungen mit Querschnittsänderungen. In: Vdi-Wärmeatlas, editors. Verein Deutscher Ingenieure VDI - Gesellschaft Verfahrenstechnik und Chemieingenieurwesen (GVC). Berlin Heidelberg: Springer-Verlag; 2006. Lac 1-8.
9. Kast W. Druckverlust in durchströmten Röhren. In: Vdi-Wärmeatlas, editors. Verein Deutscher Ingenieure VDI - Gesellschaft Verfahrenstechnik und Chemieingenieurwesen (GVC). Berlin Heidelberg: Springer-Verlag; 2006. Lab 1-4.
10. Rissler J, Jørgensen C, Rye Hansen M, Hansen NA. Evaluation of the injection force dynamics of a modified prefilled insulin pen. *Expert Opin Pharmacother*. 2008;9(13):2217-22.
11. Asakura T, Seino H, Kageyama M, Yohkoh N. Evaluation of injection force of three insulin delivery pens. *Expert Opin Pharmacother*. 2009;10(9):1389-93.
12. Papanas N, Maltezos E. The diabetic hand: a forgotten complication? *J Diabetes Complications*. 2010;24(3):154-62.
13. Carter J, Roberts A. Usability of a pre-filled insulin injection device in a 3-month observational survey of everyday clinical practice in Australia. *Curr Med Res Opin*. 2008;24(10):2741-9.
14. Haak T, Edelman S, Walter C, Lecointre B, Spollett G. Comparison of usability and patient preference for the new disposable insulin device Solostar versus Flexpen, lilly disposable pen, and a prototype pen: an open-label study. *Clin Ther*. 2007;29(4):650-60.