Patient satisfaction with the BETACONNECTTM autoinjector for interferon beta-1b

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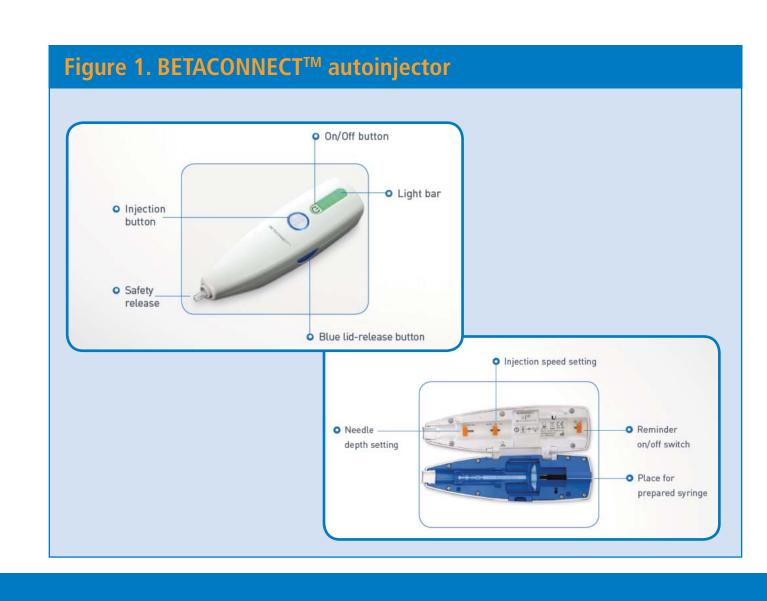
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- Treatment of multiple sclerosis (MS) requires long-term use of disease-modifying therapies (DMTs) for patients to manage their disease
- Many of the currently-approved DMTs require patients to perform regular self-injections, which can be associated with injection-site reactions or pain
- Interferon beta-1b is an injectable DMT with a favorable long-term safety and efficacy profile for patients with MS²⁻⁴
- Use of an autoinjector may reduce the occurrence or severity of some of these adverse events and increase patient satisfaction and overall quality of life^{5,6}
- Adherence with treatment may also be improved when patients use autoinjectors 7
- BETACONNECTTM is a new electronic autoinjector for interferon beta-1b that can automatically collect data such as injection date and time, injection depth, injection speed, and injection volume as patients perform injections (Figure 1)
- BETACONNECT™ has visual and auditory indicators of battery status, safety release, and injection completion, as well as visual and audible injection reminders and adjustable injection speed and depth
- The device can also transfer the collected data to an optional mobile phone application/computer program and a navigator application to enhance communication between patients and health care professionals and to help patients maintain adherence with therapy

bjective

 To assess satisfaction with BETACONNECT™ among patients using the autoinjector to administer interferon beta-1b



- Two surveys were conducted in Germany to evaluate satisfaction among patients on interferon beta-1b (Betaferon®) using BETACONNECT™ (Table 1):
- Survey 1 was conducted to evaluate overall satisfaction with BETACONNECT™ and the importance of its features
- Survey 2 was conducted to further validate the satisfaction levels observed in **Survey 1** and gain a deeper understanding of the impact of BETACONNECT™ on the patient's injection experience

Table 1: Design of the 2 surveys Survey 1 Survey 2 Patients in Germany using BETACONNECT™ were Patients in Germany using BETACONNECT[™] to All patients gave previous written consent to Additionally, patients who signed prior written consent to be contacted were sent invitations

• Recruitment approaches identified unique respondents for Survey 1 and Survey 2 to avoid overlap

Major inclusion criteria

 Participation in the BETAPLUS® program and use of the BETACONNECT™ device 	 Use of BETACONNECT[™] for ≥2 weeks, use of interferon beta-1b for ≥6 months, and performance of self-injections the majority of the time
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Survey and data collection • Data was collected from a structured paper survey | • Data was collected from a structured online survey

13-question survey

- Overall assessment of BETACONNECT™ likelihood to recommend it, and helpfulne of the device was gauged on a 6-point scale
- from 1 (most positive) to 6 (most negative)
- Importance of BETACONNECT[™] features gauged on a 4-point scale from 1 (very important) to 4 (unimportant)

• 15-minute survey - Questions on intuitiveness, ease of use, and impact on the patient experience gauged or a 6-point scale ranging from Strongly Disagree to Strongly Agree - An open-ended question on reasons for

was included

satisfaction with BETACONNECT™

Respondents

1,365 respondents completed **Survey 1**, and 118 respondents completed Survey 2 (Table 2)

Table 2. Demographic and past injection experience of patients with MS participating in each survey

		Survey 1 (n=1,365)	Survey 2 (n=118)
Female, %		69	72
Age, %	<19 years	2	0
	20-29 years	7	7
	30-39 years	18	18
	40-49 years	30	37
	50-59 years	32	29
	≥60 years	9	9
	No answer	3	0
Previous injection method(s), % ^a	BETACOMFORT®	59	56
	BETAJECT® Comfort	23	39
	BETAJECT® Lite	3	14
	Manual, without autoinjector	4	4
	No answer	11	3
Length of time using BETACONNECT™, %	<1 month	11	31
	>1 month	86	69
	No answer	3	0

^aSurvey 1 captured only the patient's injection method prior to using BETACONNECT™, while Survey 2 recorded any previous injection method the patient had used (multiple selections were allowed).

Results (cont)

Overall Impressions of BETACONNECTTM

- Overall impressions of BETACONNECTTM in **Survey 1** (Figure 2) and Survey 2 (Figure 3) were positive
- The majority of respondents preferred BETACONNECTTM over their previous autoinjector, felt it was easy to use, and were confident when performing an injection with it
- Patients were very likely to recommend BETACONNECTTM to a friend

Figure 2. Overall impressions of the BETACONNECT™ autoiniector

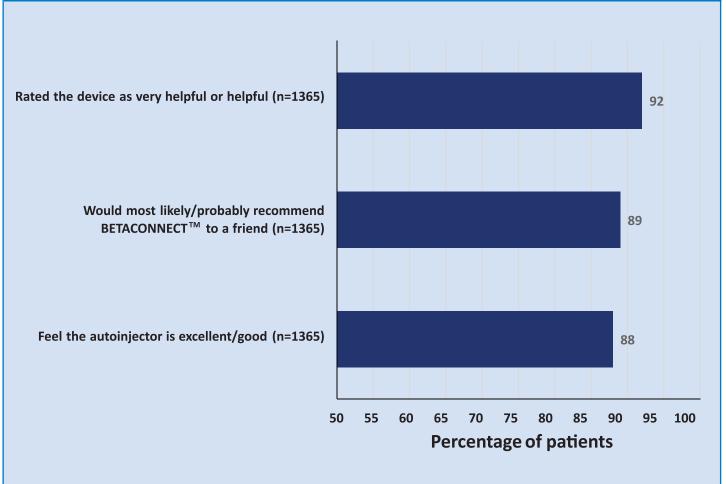
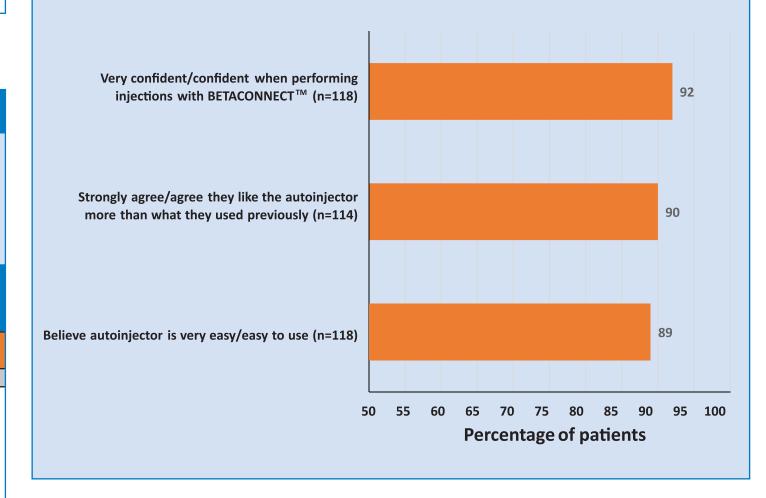


Figure 3. Overall impressions of the BETACONNECT™ autoinjector



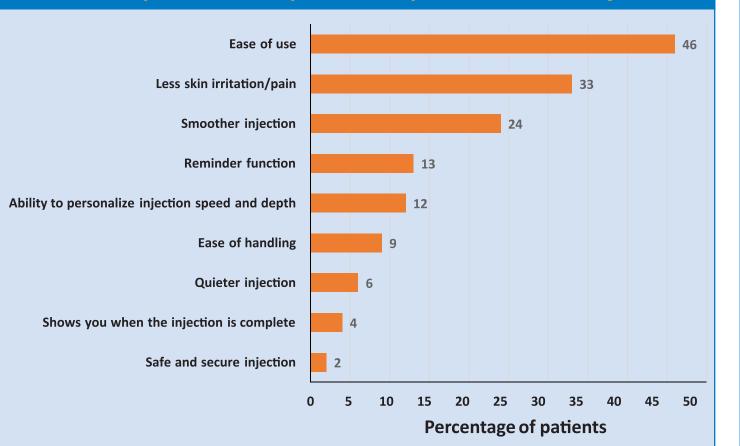
Survey 1 was supported by Bayer Vital GmbH, Leverkusen, Germany. Survey 2 was supported by Bayer HealthCare Pharmaceuticals, Inc, Whippany, NJ, United States.

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Reasons for Satisfaction with BETACONNECT™

- Unaided, patients in Survey 2 said the primary drivers for satisfaction with BETACONNECTTM were its ease of use, reduction of irritation and pain at the injection site, and smoother injections (Figure 4)
- Some patients also appreciated the reminder function, ability to personalize injection speed and depth ease of handling, and quieter injections

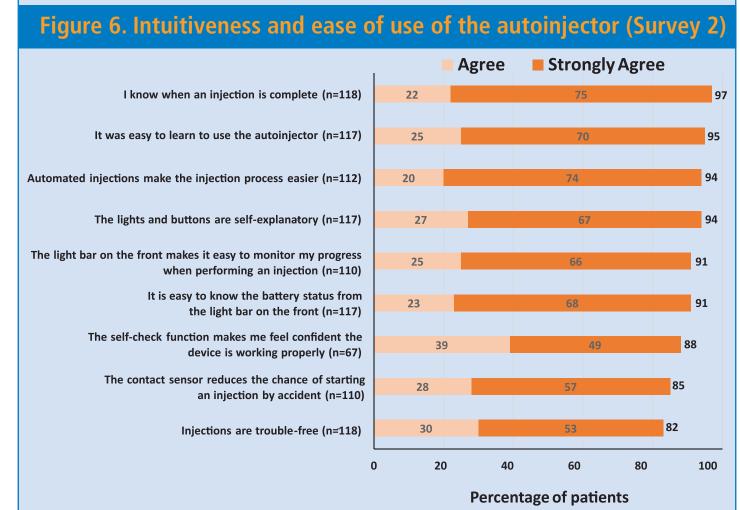
igure 4. Primary reasons for satisfaction with BETACONNECT" unaided responses to an open-ended question from Survey 2, n=



Most patients in Survey 2 agreed BETACONNECT™ was easy to learn to use and felt the lights and buttons were self-explanatory (Figure 6)

Intuitiveness and Ease of Use

The majority agreed it was easy to know when an injection was completed and that the safety release reduced the chance of starting an injection by accident



Participants who answered "don't know" or did not answer were excluded from the tally for that question

Assessment of Features

- Majority of features were rated as either "very important" or "important" by most respondents in **Survey 1** (**Figure 5**)
- In particular, patients considered the adjustable injection depth and speed, safety release, optical and acoustic signals, and rechargeable battery to be among the most important

igure 5. Importance of features from Survey 1 (n=1365)

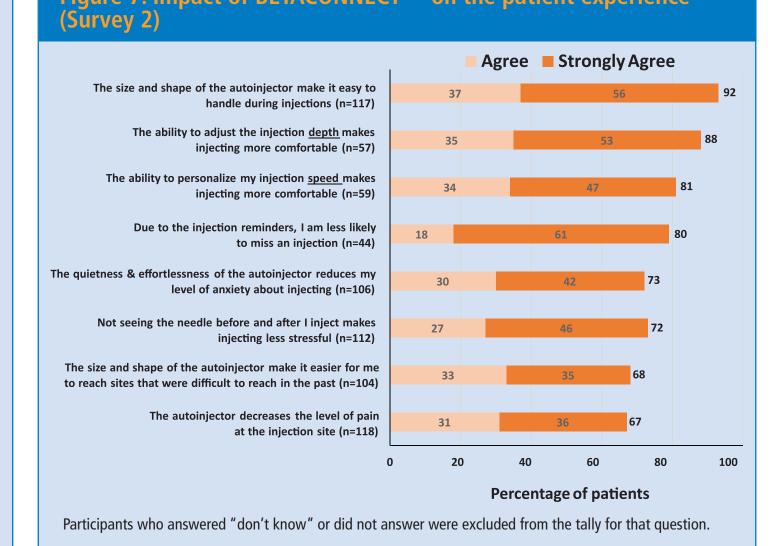
■ Important ■ Very important LED display of rechargeable battery LED display of injection process

Percentage of patients

Impact on the Patient Experience

- Participants from Survey 2 largely agreed BETACONNECT™ had a favorable impact on their injection experience including comfort of injections, ease of handling, the ability to reach injection sites, and reduced anxiety/ pain (Figure 7)
- Most patients using the reminder function said they were less likely to miss an injection

Figure 7. Impact of BETACONNECTTM on the patient experience



- The results from the two complementary surveys confirm the preference for BETACONNECT™ over the previous device and suggest BETACONNECT™ may help increase satisfaction with treatment among patients on interferon beta-1b therapy
- Overall patient satisfaction with the BETACONNECT™ autoinjector was high, driven by its ease of use, reduction of irritation and pain at the injection site, and smoother injections
- Patients believed the new features of the autoinjector were important and stated that the lights and buttons helped guide the injection process
- Participants largely agreed BETACONNECT™ improved the injection experience, including the comfort of injections, ease of handling, the ability to reach injection sites, and reduced anxiety/pain
- Most patients using the reminder function stated they were less likely to miss an injection, suggesting the autoinjector may increase adherence of patients on interferon beta-1b therapy
- Future publications will describe the impact of the autoinjector's unique communication features on adherence and information sharing between patients and health care professionals

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- IW is a salaried employee of Bayer Vital GmbH.
- JV is a salaried employee of Bayer Vital GmbH.
- AS is a student apprentice of Bayer Vital GmbH
- TS is a salaried employee of Bayer Vital GmbH.
- NP is a salaried employee of Vitartis Medizin-Service GmbH.
- APR has served as a consultant/paid advisory board member for Acorda Therapeutics; EMD Serono, Inc; Genzyme Corporation; Novartis; and Teva. She also reports receiving lecture fees from Acorda Therapeutics; Biogen Idec; EMD Serono, Inc; Novartis; Pfizer, Inc; and Teva.