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Please note that with autoISF you are in an early-dev. environment,

where the user interface is **not optimized for safety** of users who stray away from intended ways to use. Good safety features exist, but these are only as good as the development-oriented user understands and implements them. This is not a medical product, refer to disclaimer in <u>section 0</u>

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4.1 Getting started

- 4.1.1 Reminder of pre-requisites
- 4.1.2 autoISF factors overview in typical glucose curve
- 4.1.3 Getting ready to set your autoISF_weights
 - 4.2 Initial bg rise: bgAccel_ISF
- 4.2.1 Mimicking a HCL bolus
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- 20 4.2.7 Summary
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- 22 4.3 Strong bg rise: pp ISF
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- 4.5 Sluggish rise into a bg plateau, or late plateau
 - ing at high bg: dura ISF and bg ISF
- 30 4.5.1 dura ISF for sluggish rise into a bg plateau
- 31 4.5.2 dura_ISF for late/high bg plateaus
 - 4.5.3 "One size fits all" -dura_ISF
- 33 4.5.4 Options to limit iob from dura ISF
- 34 4.5.5 How dura_ISF works
- 35 4.5.6 Set your dura ISF
- 36 4.5.7 Set your bg ISF
- 37 4.5.8 "Quality control" on your tuning
- 38 4.5.9 How "UAM" concludes insulinRequ.
- 39 4.6 Tuning your initial settings
- 40 4.7 Covering more complex scenarios
- 41 4.8 Profile helper

Available related case studies:

Case study 4.1: Pizza

Case study 4.2: Low carb meal

Case study 4.3: Hands-off FCL on Xmas

43	4.1 Getting started
44	
45	Caution: This entire e-book is about Full Closed Looping (FCL).
46	In case you intend to work with giving boli, many suggestions made - notably in this
47	section 4, and in section 2 – should not be followed. Y
48	You should then primarily use the autoISF Quick Guide (from https://github.com/ga-
49	zelle/autoISF), and do extra research, on your own data. (Look at the chart in section
50	4.1.2 your bolus very much would change things there!).
51	If you shy away, for now, from FCL, please have a look into sections discussing methods
52	with "Meal Announcement", section 07, and section 13.3-
53	
54	4.1.1 Reminder of pre-requisites
55	
56	This section 4. is about the core FCL aspects of autoISF. Before doing anything with this section,
57	please make sure you have studied the preceding $\underline{\text{sections 1}}$ and $\underline{\text{2}}$ on the general pre-requisites for
58	FCL and the developers "Quick guide" (see <u>section 3)</u> on the principal workings of autoISF.
59	Core points are briefly summarized below.
60	
61	Start with proper "safety" settings
62	
63	Before you start tuning your autoISF for FCL, make sure you have appropriately:
64	• widened the SMB size restrictions (section 2.1),
65	• elevated the max allowed ISF amplification via your set autoISFmax (section 2.2)
66	Both of these points are extremely important: If you set (or keep in place) narrow restrictions,
67	this will not allow to see effects from a more aggressively tuned ISF. Even worse, it would
68	cover-up too aggressive settings (e.g. on theISF_weights that we get to in a moment), and
69	invariably make your loop bounce against the restriction(s).
70	This could even work fine, if your meal spectrum isn't broad: If, in your HCL, the same bolus
71	size pretty much fitted all your meals, it could now, in FCL, be replaced by rushing, with super-
72	aggressively modulated ISFs, into the set restrictions, to produce - with only a brief delay – the
73	required iob that would be about equivalent to what you formerly had bolussed in your HCL.
74	
75	A system that is really fit for the variance we all like to enjoy in our daily lives, though, would
76	be characterized by "tolerating" pretty wide open safety restrictions*), while having cautiously
77	calibrated other, notably ISF modulating, parameters (as described in $\underbrace{\text{sections } 4.2 - 4.5}$).
78 79	*) Still, for safety (as also suggested in section 2.1 and 2.2), start your tuning on a middle ground, and only
1 1	gradually widen SMB size and autoISFmax during your initial tuning.

80	Also make sure you have
81	• set your iobTH% (refer to section 2.4 and if available 4.8)
82	
83	Furthermore, in your early test phase, it is recommended to:
84	• Run the system as dummy, not connected to your body (or, on own risk, connect only as long
85	as you watch closely)
86	• In AAPS preferences, switch your autoISF FCL (= autoISF/"Enable adaptation of ISF to
87	glucose behavior") ON only during daytime hours of a meal, e.g. 11-18h, for fully automatic
88	"full closed loop" management of lunches.
89	You can do this switching manually at 11 h and 18 h every day, or set up an
90	Automation that does that (see $\underline{\text{section 3.4}}$).
91	Take typical but not extreme meals. Omit sweet drinks, or drink only slowly. You are going for a
92	"good enough" compromise, that works with your range of usual meals.
93	
94	It is wasted time to do a lot of iterations to "optimize" settings based on just 1 type of meal.
95	See case study 8.2
96	
97	Occasionally, watch the time-pattern of bg, iob (SMBs given), and insulin activity after meal start.
98	Aside from serious "mathematical" attempts to tune settings based on data from the SMB tab (or
99	the Emulator, section 10), just watching the curves develop on your AAPS main screen can, over
100	time, give you "a feel" what settings, and eating behaviors, are benign or detrimental to good %TIR
101	performance.
102	
103	Importance of proper profile ISFs.
104	
105	Starters on autoISF FCL who are coming from using HCL with dynamic ISF must be aware of the
106	following: It is absolutely essential to build your FCL on a properly set profile ISFs (likely a
107	circadian pattern over 24 hrs).
108	
109	It may not apply to you, but many dynamicISF users did never bother to determine their ISFs that
110	would maximize their HCL performance, but employ dynamicISF so to speak for going
111112	"dynamically" through a wide range of possible ISFs, until eventually hitting a sweet spot, and the whole thing works better than before, with what they had <i>used</i> as a profile ISF (often only one, e.g.
113	coming from Autotune).
114	
•	

115	The following is important to understand, as it leads straight into the core idea behind FCL with
116	autoISF, too: It is a good idea to establish a well-running hybrid closed loop with set (non-dynamic)
117	ISF (set in profile for each hour of the day). That ISF must be aggressive enough that it gets you
118	down from a high around 200 mg/dl to target. That is roughly also the way you experimentally
119	determined it (so I hope. See https://github.com/bernie4375/HCL-Meal-MgtISF-and-IC-
120	settings/blob/HCLsettings-main-repo-(pdf)/ISF%20determination V.3.33.pdf).

- Using *that strong* value also *at lower bg*, (on the way "up", after meal start), is very positive: We do *not* want to have a *softer* acting loop when at *lower* bg (which is what dynamicISF tends to do!). autoISF will, in contrast, temporarily sharpen your ISF when, at low bg, acceleration is detected..
- On the way down from peak, towards glucose target, a somewhat too strong ISF should not hurt because much of the time your loop (well supplied with insulin before, "on the way up") is zero temping, or at least has only a small gap to correct, from predicted bg to target bg.
- You have no business to be much above 200 mg/dl where an even stronger ISF may or may not help. It sure does not help at an occlusion which is about the only reason to see super high values as an experienced looper.

Pegging ISF strength to bg level therefore **does not make sense in FCL**. You will use the autoISF toolbox to get strongest ISF **at low** but beginning-to-rise bg,

Note: There are very much refined versions of dynamicISF that can have beneficial applications, notably in HCL ...And, yes, I know, bg levels can also correlate with insulin sensitivity. But let us not get into "chicken or egg type" discussions.

Rather, focus on doing a good tuning job, and use superior approaches to account for sensitivity changes in a more pro-active manner (before running into sky-high bg (or into hypos)):

Going to autoISF FCL, you absolutely must anchor on the proper profile_ISF.

The profile is not "set in stone", though. To use above terminology again: **Pegging ISF strength to** your current **insulin sensitivity** – very much like you had done all along in HCL - **does make** sense in your FCL...

(...and the fact that autoISF afterwards "anyways" often strongly modifies ISF is not a reasonable counter-argument).

There are fully automated, as well as manual ways for sensitivity adaptations of the profile ISF:

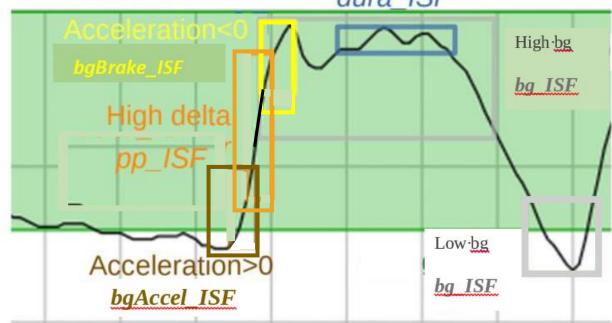
Profile ISFs can get **fully automatically adapted**, e.g. by Autosens, or by the Activity Monitor, which in autoISF we rather use (section 6.5).

153 Which of your basic related settings (in AAPS/Preferences) produce exactly which adaptation can be seen right in the top lines of your SMB tab, at each loop decision. Likewise, it can be retrieved 154 155 later in logfile analysis (see Emulator, section 10) 156 Furthermore, when using autoISF you can – as you did in the past, e.g. around exercise. or in 157 158 times of illness – temporarily **manually modify** your profile ISFs 159 160 Also these effects are quantified in SMB tab and logfiles *). 161 *) Furthermore, the results from autoISF are explained in the SMB tab, and multiplied with (original or adjusted) 162 profile ISF to result in the ISF (called "sens") used in the current insulinRequired calculation 163 164 All three top buttons in AAPS (%profile switch, exercise and TT) can be freely used to adapt to 165 changes in sensitivity/resistance, turning into a yellow color to alert you to this. (More about your 166 "FCL cockpit" see section 5.2.2.). 167 168 For a start, please spend a couple of days (if not weeks) to get your key autoISF related settings 169 right, strictly on/for days with your normal insulin sensitivity. This is what this section 4 is 170 about. 171 172 Importance of starting from a well-performing Hybrid Closed Loop 173 174 A satisfying performance in Hybrid Closed Loop mode is a pre-requisite. Expect to reproduce 175 about the same %TIR also in your FCL, but with less daily interaction, once established. Note that this refers to prior use of "vanilla" software, without fancy "dynamic add-ons" (such as: 176 177 Autotune determined factors, dynamicISF etc), that may have introduced bias into the profile 178 settings you bring with you into FCL now. 179 180 To reach a satisfying performance you must start from a hybrid closed loop in which you did 181 master your meal management well using the oref(1) algo SMB+UAM. 182 This is a pre-requisite to be able to forget it ... - because the initial tuning that we now turn to 183 demands, that you analyze your prior best practice as your blueprint to find appropriate settings 184 and "teach" your FCL to come up with the necessary iob. 185 186 This is the main subject of this section 4 (finding settings for automatic meal management). 187 188 189 190

191 Do not copy settings from other FCL loopers 192 193 When setting your parameters, don't use any given numerical example (not even as "a starting 194 point"). Instead, anchor on data from your successful Hybrid Closed Loop! 195 196 Most examples given in this paper are from an adult diabetic (Lyumjev, G6) whose insulin sensitivity 197 can be characterized as follows: approximately 37 U TDD, thereof 13 U profile basal, at about 200g 198 daily carbs from mainly lunch and dinner; no couch snacks or sweet drinks. The user also 199 participates in multiple instances of daily moderate exercise such as dog walking, biking and 200 gardening. In Hybrid Closed Loop, a typical meal bolus was 8 U that was sometimes reduced such 201 as when activity followed the meal. 202 After seeing some more inputs from a variety of users we might put together a profile helper for 203 some rough orientation, and for plausibility cross-checking, in section 4.8 204 205 Importance of going step-by-step 206 207 Section 5 will explore avenues to manage "disturbances", i.e. time blocks or situations that might 208 demand enhanced or reduced loop aggressiveness. 209 Section 6 will focus on the exercise mode, and the activity monitor. 210 In case you have a strong interest in the Activity monitor (section 6.6), you can start with 211 calibrating that, and have it run already in the weeks when you go through sections 4 and 5. 212 In case you use an EatingSoonTT at meal start (the author recommends to try without), note that any 213 active TT shuts activity monitor automatically off while that TT is active. 214 215 Resist the temptation to make use of the tools presented in sections 5 and 6 too early. 216 On your first setting-up and tuning attempt, it is strongly recommended that you not "play 217 around" with all ultimately available features, but stick to the sequence of steps to take. 218 219 Yes, "playing around" with the many extra buttons often will help find an improvement. But you 220 likely create an instable FCL that, already at fairly standard situations, uses up some of your FCL's 221 principal capacity to correct for disturbances. This limits what will be left to manage extreme 222 situations. 223 224 Caution: Once you created a maze of little errors and counter-strategies/counter-errors, it will be 225 nearly impossible to find your way out of this mess, towards better settings, at any later point of time. 226 227 Note that it is principally not easy to conclude on suitability of tuning: 228 AutoISF comes with very many (currently 18) extra parameters, and even when employing the 229 emulator (sections 10 and 11), it is quite hard to analyze their interaction.

230 One principal reason why things are difficult to analyze is, that you really can only analyze one 231 decision, and that will put you on another bg curve. So, you can never see the full effect, along more 232 than half an hour or so, that any change would really result in. 233 234 Understandably, many loopers rather "move forward" to an over-patch for identified problems, and 235 not bother with a more "puristic" step-by-step approach to do things right from the ground up. 236 Aware of above sketched conundrum, the AAPS autoISF developers offer the ultimate tool to 237 investigate "what-if", regarding a setting change you may contemplate: A nice lady voice on your 238 smartphone can tell you, at each loop decision, where your contemplated change would make a 239 difference (in SMB size). This offers an opportunity to watch closely, with or without implementing 240 that change. (It is always your spontaneous choice, whether you want to "follow the lady's 241 suggestion and manually add to the SMB, as suggested). More see in Section 11.4 242 But, we are getting ahead too far here. You first must find a starting point for key settings, which works 243 reasonably for not too-challenging meals in your personal spectrum. 244 Before getting into this, let's first have a look on how autoISF basically works. (More see in Quick guide by 245 the developer, referenced in section 3.2; or directly at https://github.com/ga-zelle/autoISF). 246 247 4.1.2 autoISF factors overview in typical glucose chart 248 249 The core challenge of your UAM Full Closed Loop is to recognize a meal start from the glucose 250 trend, and ramping up iob. 251 252 When setting up your autoISF Full Closed Loop, you must set several ISF_weight parameters 253 in AAPS Preferences/OpenAPS SMB/autoISF settings. 254 255 They relate to different stages of the typical glucose curve after starting a meal:

Plateau above target dura ISF



Note: **bg_ISF** is not used much in FCL, as it is rather late to act on high (or low) bg level that developed. But, feel free to experiment, e.g. in case you have indications, in your data, that in the past dynamicISF was useful to manage bg extremes in some situations.

The core advantage of using autoISF with oref(1) SMB+UAM (in FCL as well as in hybrid closed loop) is that it manages the glucose curve it sees developing, **no matter what the underlying reason** is.

42 potential factors were identified (see: https://github.com/bernie4375/HCL-Meal-Mgt.-ISF-and-IC-settings/blob/FCL-w/autoISF/42%20factors%20influence%20bg.pdf), so, no wonder, that loopers who meticulously input their carbs will often *not* see the expected result.

In case you like to learn more about the theoretical background for the autoISF development, here some core aspects the key developer shared in a DIY forum:

Theory behind autoISF

gazelle, Aug. 2024

I have recently managed to work through the large pile of older (German) diabetes journals. I've now reached issue 5/2024 and was amazed by the article "How insulin is produced and works in the body".

According to the article, there are essentially 2 phases of release:

- Phase (1) lasts only a few minutes and releases a large dose from the beta cell's supply when glucose rises.
- (2) After that, insulin is newly produced and released evenly over a period of hours.

These two phases remind me very much of initial acce_ISF and later dura_ISF in autoISF.

This may explain why it works so well for many users. I had no idea about this process in the body when I developed both effects.

- acce_ISF was based on Newton's 2nd law of mechanics, i.e. inertial mass * acceleration = force.
 For me, this meant that when the FC is accelerated, a "sweet force" is acting.
 To counteract this, the acceleration must be counteracted.
 Strengthening the ISF, i.e. increasing resistance, corresponds to an increase in inertial mass, so to speak.
- dura_ISF was an analogy to the PID controller in technology:
 P stands for proportional control, in this case the deviation of the FC from the target value;
 I stands for integral of the deviation, in this case the sum of the deviation from the target;
 D stands for differential control, in this case the delta of the BG.

288 The components for P and D are contained in the regular AAPS, but the I component was missing. So I added it, as a rough 289 approximation, using dura_ISF. 290 291 292 293 294 295 I find it very interesting that both solutions came about from analogies to technology/mechanics without direct knowledge of hormone metabolism. I have always been interested in using analogies to transfer solutions to other fields. Translated with DeepL.com (free version) Remark: This just broadbrush sketches the general idea. Lots of dev work had unfortunately to go into details, like how to deal with everyday imperfections of our CGM systems, and still come up with meaningful mathematical (parabola fit) 296 297 earliest-possible detection of "real" bg rises. 298 4.1.3 Getting ready to set your autoISF weights 299 300 Before you progress, make sure you studied the flowcharts in section 3 that describe how autoISF 301 calculates the effective(ly used) ISF. 302 303 Warning: Any bolus you "sneak in" will severely distort the glucose curve. That could 304 render your tuning of weights (see below) useless, and could make your loop act in 305 unpredictable ways (potentially also dangerous, however, your set iobTH (section 2.4) should 306 help here, too). 307 308 In case you feel tempted to use boli, be ready for some own extra research, and refer to 309 section 7. 310 After doing the prep work as outlined in section 2 you now get to calibrate your FCL to your 311 312 normal meal spectrum by initially setting and tuning the various ISF weights, that 313 dynamically change with bg curve characteristics as sketched in the chart on the previous page. 314 315 Please stay away from extremes (regarding both, meals and exercise) when you go through 316 this section 4. It is about getting a first roughly right set of settings, as a basis. 317 318 Researching your standard meal patterns, and finding settings for the various -ISF_weights 319 is the core job in setting up your autoISF FCL. 320 Depending how varied your diet and general lifestyle are (and your expectation of %TIR 321 you like to reach), this could be the main job at hand. However, there is much more you 322 could do later, and that will be outlined in later sections 5 and 6. 323 324 Consult sometimes your SMB tab, to see how the applied effective ISF (named sens there) is 325 calculated. (Example given in section 5.4.5). 326

327	4.2 Meal detection and managing the initial bg rise: bgAccel_ISF
328	
329 330	4.2.1 Mimicking a HCL bolus in FCL using bgAccel_ISF
331	When looping without carb inputs and without giving a bolus ourselves, the first crucial setting is to
332	set the bgAccel_ISF_weight so that SMBs are requested immediately when the loop detects an
333	acceleration in your blood glucose (bg) that is starting to rise.
334	
335	Ideally within about 20 minutes after acceleration detection, which would be the first up to 4
336	SMBs, as much iob should automatically be supplied as we would have given with our
337	bolus in hybrid closed loop.
338	
339	As the biggest principal challenge for the FCL is big high/fast carb meals (from within your
340	personal "spectrum"), we start with a focus to get sufficiently big SMBs going for those.
341	
342	Note, though, that in a low carb meal scenario, the first 4 SMBs would have to automatically result
343 344	much smaller (which, after careful tuning, is possible with the same parameter settings, see e.g. case studies 4.2 vs 4.3).
345	case studies 4.2 vs 4.3).
346	Rule of thumb: Two of the first three SMBs each (in this test based on a big meal) should be about
347	$\frac{1}{4}$ to 1/3 the size of a bolus in your HCL "career" (for a similar meal).
348	Going over 1/3 could be problematic
349	• if your diet contains occasional low carb (or brief snack ing), it is not helpful if your
350	settings make your loop invariably "bounce" over your iobTH (and then you would
351	need extra snacks to balance the auto-generated iob, to prevent hypos),
352	• also if your CGM quality is sometimes unreliable, and might produce an artefact
353	that could be mistaken for a meal start.
354	Be vigilant about this topic! And please do not choose the supposedly easy way, to just set safety
355	restrictions (allowed max SMB size, or autoISFmax) so low, that your loop never can exceed 1/3.
356	Try to really tune the _ISF_weights appropriately. (Only that way, your loop can "accommodate" the
357	entire meal spectrum, and also states of adapted general insulin sensitivity).
358	
359	4.2.2 Widened safety restrictions
360	
361	Already when tuning the bgAccel_ISF_weight it can become evident that safety restrictions (as
362	discussed in section 2) must be widened further:

363 Especially if your *profile basal* rate is very small, the **smb_max_range_extention** and/or 364 the autoISF max "must" often be increased further. 365 • Pay attention also to the **iobTH**% and, potentially, iobMAX 366 Note that the smb delivery ratio "only" portions the insulinReq differently over the next 15 367 minutes (see also section 2.3), and therefore is **not** a prime tuning parameter. 368 In the end you should **not set these safety limits too tight,** so "nudging" aggressiveness by 369 another 10 or 20% from your cockpit, later, will not bounce into restrictions. 370 371 On the other hand, setting **narrower** restrictions for max allowed SMB size can also become 372 necessary: 373 Poorer CGM quality demands narrower restrictions for safety reasons. 374 If you use a 1-minute CGM, please observe section 1.4.2 375 376 4.2.3 Start value for your bgAccel ISF tuning 377 378 bgAccel ISF weight is set default to zero in AAPS Preferences/SMB/autoISF. 379 **To start**, I would try 0.05 or **max 0.1**, and keep trying in max 0.05 steps. Soon move to 0.02 steps. 380 From my (very limited) overview, many use around 0.2, and possibly even higher if their hourly 381 basal rate is 0.1U or lower. (Consult section 4.8 when available). Do not be tempted to rush this 382 setting by using large jumps in adjustments. 383 To monitor what is happening, and start tuning, in search of appropriate settings, you must keep 384 (real-time) track of how autoISF uses your set bgAccel ISF weight: 385 386 To do this in the **SMB tab** is possible but not very practical. You would end up making a lot 387 of screenshots (quickly in the crucial minutes after a SMB was given, or when you thought it 388 should be given), for later analysis. 389 The superior method is to just copy logfiles from your phone/internal 390 memory/AAPS/logs ... 391 o all zip files there 392 o look up how many days of data are covered there on a rolling basis, and copy out 393 onto your PC (see section 10.1.1) before the older ones get forever lost 394 ... and analyze them at your convenience later, using the emulator (see section 10; used 395 e.g. in last pages of case study 4.2). 396 • Some emulator-based analysis is also possible within AAPS on your phone (section-11).

```
397
       In any case, it is worth the effort to tune the bgAccel_ISF_weight in such a way that high glucose
398
      increases are already nipped in the bud, so to speak.
399
400
       To summarize: In FCL, the first 3 or 4 SMBs should not be much delayed, and amount to similar
401
      iob like your "former boli in HCL".
402
       Depending on details about the carb absorption characteristics of your meal, and the performance
403
       of your CGM, also pp_ISF (see 4.3) might be a fairly early contributor to getting iob up.
404
405
      4.2.4 How changing the weights influences the resulting calculated insulinRequired
406
       (You can skip reading this section, unless you want to know quantitatively how things work).
407
408
      The developers' documentation (Quick Guide) https://github.com/ga-
409
       zelle/autoISF/blob/A3.2.0.4 ai3.0.1/autoISF3.0.1 Quick Guide.pdf gives the following equation:
410
411
      accelSF is the factor by which autoISF wants to sharpen the profile ISF in a certain
412
       situation of bg acceleration:
                 acce ISF is calculated by
                                                                                               (eq.1)
               acce_ISF = 1 + acce_weight * fit_share * cap_weight * acceleration
                            a measure of fit quality, i.e. 0% if unacceptable up to 100% if perfect;
        where fit_share
                            is 0.5 below target and 1.0 otherwise;
               cap weight
                           is bgAccel_ISF_weight for acceleration away from target, i.e. mostly positive
               acce_weight
                            or bgBrake_ISF_weight for acceleration towards target, i.e. mostly negative
413
414
              fit_share should be close to 1.0 (if good CGM quality.
415
              cap weight is 1.0 for bg>target
                           Having a EatingSoonTT at first acceleration can help avoid the factor iF getting cut in half!
416
417
              profile ISF / acce ISF = effectively used ISF (sens)
418
               (...if the acce influence dominates and is used as effective ISF. Else, see flowcharts in Quick Guide)
419
420
      When looking at the same acceleration moment, we can combine all three last factors of eq.1 into a term
421
422
      For estimating the effect from using another ISF weight, we then have two (eg.1), with two unknowns, iF
423
       ("cancelling out"), and the sought acce_ISF (for new _weight).
424
425
               Example: Your profile ISF is 40 mg/dl/U. Using bgAccel ISF weight of 0.2
              you saw effectively used ISF of \frac{30.3}{1.3} mg/dl/U (box C13 in table below) = 40 / 1.32
426
427
              => factor for acce ISF = 1.32.
428
              For an intended correction by – 10 mg/dl the insulinRequired would calculate to 10 / 30. 3 =
429
              0.330 U.
```

4	Α	В	С	D	E	F	G	Н
1	Profil ISF (mg/dl/U)	40		bgAccel_ISF	tuning -			
2	InsReq (U) @ iF=0	1		see FCL e-Bo	ook section 4.2.4			
3	bgAccel_ISF_weight:	0,2			0,1			InsReq effect
4	iF	acce_ISF_old	ISF_old	InsReq_old	acce_ISF_new	ISF_new	InsReq_new	
5	0	1	40	1	1	40	1	0%
6	0,2	1,04	38,5	1,04	1,02	39,2	1,02	-2%
7	0,4	1,08	37,0	1,08	1,04	38,5	1,04	-4%
8	0,6	1,12	35,7	1,12	1,06	37,7	1,06	-5%
9	0,8	1,16	34,5	1,16	1,08	37,0	1,08	-7%
10	1	1,2	33,3	1,2	1,1	36,4	1,1	-8%
11	1,2	1,24	32,3	1,24	1,12	35,7	1,12	-10%
12	1,4	1,28	31,3	1,28	1,14	35,1	1,14	-11%
13	1,6	1,32	30,3	1,32	1,16	34,5	1,16	-12%
14	1,8	1,36	29,4	1,36	1,18	33,9	1,18	-13%
15	2	1,4	28,6	1,4	1,2	33,3	1,2	-14%
16	2,2	1,44	27,8	1,44	1,22	32,8	1,22	-15%
17	2,4	1,48	27,0	1,48	1,24	32,3	1,24	-16%
18	2,6	1,52	26,3	1,52	1,26	31,7	1,26	-17%
19	2,8	1,56	25,6	1,56	1,28	31,3	1,28	-18%
20	3	1,6	25,0	1,6	1,3	30,8	1,3	-19%
21	3,2	1,64	24,4	1,64	1,32	30,3	1,32	-20%
22	3,4	1,68	23,8	1,68	1,34	29,9	1,34	-20%
23	3,6	1,72	23,3	1,72	1,36	29,4	1,36	-21%
24	3,8	1,76	22,7	1,76	1,38	29,0	1,38	-22%
25	4	1,8	22,2	1,8	1,4	28,6	1,4	-22%
26	4,2	1,84	21,7	1,84	1,42	28,2	1,42	-23%
27	4,4	1,88	21,3	1,88	1,44	27,8	1,44	-23%
28	4,6	1,92	20,8	1,92	1,46	27,4	1,46	-24%
29	4,8	1,96	20,4	1,96	1,48	27,0	1,48	-24%
30	5	2	20	2	1,5	26,7	1,5	-25%

31 formula (line 30) =1+\$A30*B\$3 =\$B\$1/B30 =+\$B\$2*\$B\$1/C30 =1+\$A30*E\$3 =\$B\$1/E30 =+\$B\$2*\$B\$1/F30 *=G30/D30-1
32 Note: To determine not the incremental effect between two weights (E3 vs.B3), but the acce_ISF effect
33 for the investigated _weight (E3): Enter 0 in (B3)
34 Column (H) then shows the full effect of acce_ISF with the selected weight (E3)

Now, going with a 50% reduced bgAccel_ISF_weight of 0.10:

acce_ISF = 1+ bgAccel_ISF_weight * internalFactor

before 1,32 = 1 + 0.20 * iF => 0.32 = 0.20 * iF => iF = 1.60 (see box A13)

after ? = 1 + 0.10 * iF => ? = 1 + 0.10 * 1.6 = 1.16

436 New effective ISF would be $40 / 1.16 = \frac{34.48}{40} \text{ mg/dl/U}$ (box F13).

For an intended correction by – 10 mg/dl the insulinRequired would now calculate to 10 /

To open the ...xls, go to: "Download raw file"

34.48 = 0.290 U, which is 12 % less (H13).

At higher acceleration points of the bg curve (column A, line 14 and higher), the effects would be more pronounced (column H, line 14 and higher).

An **Excel tool** is provided in e-book section 04 for this calculation. However, easier to handle

might be the:

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Simple "trial and error" approach

To get a feel for how changing the _weights influences the resulting calculated insulinRequired, it is best to start cautiously and just do 10 to max 20% steps up, and watch out for the effects.

450 Doing similar step sizes should yield about similar effects each time.

451	Never forget to look into how otherISFs play into the effective ISF (named sens in the SMB tab),
452	which overall results.
453	4.2.5 Characteristics of a well tuned-in bgAccel_ISF_weight
454	
455	Your starting point was to set the bgAccel_ISF_weight so FCL works in a rather high carb meal.
456	
457	Now you must check (and potentially fine tune) so it will not "shoot iob too high" with the first 3
458	or 4 SMBs in other meals from your spectrum:
459	
460	For meals that are in the lower (!) range of the "fast carb load " of your cluster, the
461 462	necessary insulin supply for the first two hours or so might pretty much be provided already with the first 3 or 4 SMBs
460	
463 464	The glucose curve, at such meals, begins to flatten early in this SMB phase, so a deceleration (braking) follows very soon (-> section 4.4). Clearly, the first 3 SMBs, in such
465	cases, must remain below iobTH.
705	cases, mast remain below lost in
466	Low carb meals are principally easiest for the FCL. However, you must secure that your
467	bgAccel_ISF driven first SMBs remain small. This is principally possible also with a fairly
468	aggressive bgAccel_ISF_weight set, because both acceleration and initial deltas are small
469	when eating low carb. (Regarding the detected acceleration, the stakes may be high for the
470	CGM and smoothing method you chose).
471	 A stage where moderate amounts of carb absorption and of insulin usage/need hold a
472	balance could protract – at moderate bg elevation -over hours. The dura_ISF might play a
473	bigger role, then, as e.g. in the low carb example in <u>case study 4.2</u> .
474	
475	In case you run into limitations, see next sub-chapter.
476	
477	4.2.6 Suitability for many types of meal
478	
479	For a hands-off FCL, your settings have to fit
480	• in each of your meal times
481	What helps here is that, between your daily mealtime slots, your circadian profile ISFs
482	(upon which the autoISF modulations build) automatically make a differentiation (as was
483	the case in your HCL).
484	• for the whole range of <u>your</u> meals. All this is principally possible, but:

485 486	What if you still have meals that you cannot make fit?
487 488 489 490	In extreme cases you will have to balance too high running iob with additional carbs (a late additional snack against going too low), and in the opposite case, you will have to reckon with temporarily exceeding the glucose target range, and losing some %TIR for this day.
491 492	If your meals vary very strongly, there are avenues to ease your initial tuning job, or to optimize overall resulting loop performance:
493 494 495	 Automations allow you to differentiate. For instance it is possible to apply different iobTH_percent and/or different bgAccel_ISF_weights for meals in different time windows or geo locations (details see <u>sections 3.4</u> and <u>5.1)</u>.
496 497	In case you use autoISF on the Trio or iAPS platform for i-phones, you may need to use a third party automation software, or "middleware" (! call for a case study $4.X$)
498 499	 you can pre-program custom buttons for special meal (or snack) types, with different underlying FCL settings (see "cockpit", section 5.2.2.3)
500501502	 You can modulate FCL aggressiveness manually making use of the top 3 buttons in the AAPS home screen: These turn yellow during temporary switched %profile or glucose target (section 5.2.2.2)
503504505	 Experimenting with the three above mentioned "avenues", the author found: the last point easiest to occasionally use, and the first one hardest.
506 507 508 509	 it worth investing some effort (also using the emulator a couple of times) to iterate through the typical meal spectrum a couple of times, for finding a "good enough" set ofISF_weights and other settings (like autoISFmax, iobTH% etc), and not do much extra differentiation. (More see in section 5).
510 511 512	4.2.7 Summary on tuning for the initial SMBs via bgAccel_ISF
513514515516	Early strong iob also will ease the tuning task for the subsequent phases of the meal, because there is, then, largely zero-temping (as well known from HCL-times after your administered bolus). Also, the lower and shorter lasting the glucose peak, the lesser the hypo danger from the activity tail of SMBs given <i>when</i> glucose was "stuck" high.
517 518 519	However, it is important not too super-aggressively tune bgAccel_ISF_weight up, so, regardless of the type of meal, very big SMBs invariably would result. Rather, the rough idea should be:

520 521	 SMBs driven by bgAccel_ISF: initial iob for all meals. SMB sizes vary, because accelerations and deltas vary.
522 523 524	So, at high carb meals it depends on your settings, and on the evolving bg curve, whether the first few bgAccel_ISF driven SMBs get you already up to iobTH in high carb meals, or whether this happens in the <i>overlapping</i> next stage.
525	So, looking a bit ahead to the next chapters:
526 527	• SMBs driven by pp_ISF: to the extent there is strong (near-linear) bg rise (at big meals rich in carbs) with big or small deltas, iob is now driven towards (and potentially over) iobTH.
528 529	In low carb meals this period can be extremely short, with iob remaining under iobTH (example see case study 4.2)
530	SMBs driven by bgBrake_ISF, bg_ISF, or dura_ISF:
531532533534535	Note that <i>all of these</i> can overlap with the pp_ISF stage. Consult the csv table output from the Emulator (example given at end of <u>case study 4.2</u>) as to which of the _ISF categories drives the effectively used ISF (and what change of theISF_weights would change this. Consult decision flowcharts for effective_ISF in pages 1-6 of the Quick Guide.pdf in https://github.com/ga-zelle/autoISF).
536537538539	Depending on the shape of the bg curve after the initial strong rise, and depending on insulinReq. and on iob (> iobTH?), autoISF can provide more SMBs to bring bg to target. This case applies to low carb meals. The dura_ISF is also useful to manage temporary insulin resistance often observed late in fatty meals.
540	
541	It is worth investing effort (following the sequence of steps in sections 01-04 of this FCL e-book)
542	in your initial project to establish a good set of ISF_weights for your meal spectrum. This will
543	keep interventions in daily life to a minimum.
544	Unless your lifestyle, or health and body weight change radically, this should be a <i>one-time</i> effort
545	(in your initial weeks establishing your FCL), with <i>no need</i> to fine-tune much later (see <u>section 8</u>).
546	
547	4.2.8 Note regarding acceleration "happening again" in late part of <u>dropping</u> glucose
548	(Skip, unless interested)
549	After the peak, in the late stage of falling bg, the glucose curve is like an accelerating
550	parabola again. The algorithm tries to evaluate when and at which bg level complete
551	digestion of the meal and a bg minimum will result. Insulin required to stabilize around
552	target bg is usually very small, and the adaptation of ISF in that stage relatively
553	unimportant. See in your SMB tab, how, at "already falling" bg, the ISF modulation is taken
554	back.

In version 2.2.8.2 there was a potential deficiency in situations where glucose was falling and the glucose acceleration was already positive. That meant a minimum glucose level can be extrapolated. If that happens to be less than target and expected in less than 15 minutes then there should be no strengthening of ISF as it would lower glucose even more. Therefore bgBrake_ISF_weight is used now instead of bgAccel_ISF_weight. But those situations were rare and less critical than might be expected at first sight. The reason is that in most cases the predictions ended up even below their threshold meaning SMB were disabled.

4.3 Managing strong bg rises: pp ISF

4.3.1 Main function of pp_ISF in autoISF FCL

In the later phase of acceleration and in the earlier phase of deceleration there is a more or less linear increase of bg with **high deltas**, and corresponding extra insulin need.

• With **higher carb load** meals, or meals that come with a sweet drink, the increase will be particularly strong, and (if not already driven there by bgAccel_ISF) now reach, and with the last "allowed" SMB exceed, the valid iobTH.

• With **low carb** meals, there is only a very un-pronounced (short, with weak deltas) "pp_ISF phase". (Example see end of case study 4.2).

autoISF should now "fight" this with the help of the post-prandial ISF, set via **pp_ISF_weight**, after you have set your bgAccel_ISF_weight.

4.3.2 Tuning pp_ISF_weight

To tune-in your **pp_ISF_weight**, please do this with a really high carb meal (from within your typical meal spectrum) *after* you have set a halfway suitable (not too aggressive) bgAccel_ISF_weight.

Note that if you rush into pp_ISF tuning while "still having a too aggressive bgAccel_ISF", the latter is covering up the requirement you now really want to calibrate for in pp_ISF!

So, at a meal in the upper spectrum of your carb load, carefully begin with a starting value for pp_ISF_weight of 0.005. Observe the reactions and check the SMB-tab before you increase it cautiously for the next days.

Best practice is to analyze the emulator tables (discussed in <u>section 10</u>, and example given in the pizza <u>case study 4.1</u>)

586 How changing the weight influences the resulting calculated insulinRequired 587 (You can continue 2 pages down at 4.3.3, unless you want to know quantitatively how things work). 588 589 The developers' documentation (Quick Guide, page 5) https://github.com/ga-590 zelle/autoISF/blob/A3.2.0.4 ai3.0.1/autoISF3.0.1 Quick Guide.pdf gives the following equation: $pp_ISF = 1 +$ (eq. 2)delta * pp_ISF_weight. 591 592 profile ISF / pp ISF = effectively used ISF (sens) 593 (....if the pp influence dominates, and is used as effective ISF. Else, see flowcharts in Quick Guide) 594 595 Note that the pp ISF effect only comes in when there is positive short avg delta, AND bg 596 is minimum 10 mg/dl above bg_target already. 597 Before, we need bgAccel ISF do its job! 598 Evidently, by setting a lowTT around meal start, you can influence (not the size, but) 599 how soon pp_ISF will "compete with" your bgAccel ISF (... and this can play out 600 very differently, at different kinds of meal, too.) 601 602 Tuning: When looking at the same moment (same delta), the effect from using another ISF weight, 603 can be mathematically be solved with two (eq.2): 604 605 Example: Your profile ISF is 40 mg/dl/U. Using dura ISF weight of 0.02 you saw 606 effectively used ISF of $\frac{30.3}{1.3}$ mg/dl/U (box C13 in table below) = 40 / 1.32607 => factor for pp ISF =1.32.

For an intended correction by - 10 mg/dl the insulinRequired would calculate to 10 / 30. 3 =

608

609

0.330 U.

	Α	В	С	D	E	F	G	Н
1	profile.ISF (mg/dl/U)	40		pp_ISF tunir	ng -			
2	InsReq (U) @delta=0	1		see FCL e-B				
3	pp_ISF_weight:	0,02			0,03			InsReq effect
4	delta (mg/dl)	pp_ISF_old	ISF_old	InsReq_old	pp_ISF_new	ISF_new	InsReq_new	(E3 vs.B3)
5	0	1	40	1	1	40	1	0%
6	2	1,04	38,5	1,04	1,06	37,7	1,06	2%
7	4	1,08	37,0	1,08	1,12	35,7	1,12	4%
8	6	1,12	35,7	1,12	1,18	33,9	1,18	5%
9	8	1,16	34,5	1,16	1,24	32,3	1,24	7%
10	10	1,2	33,3	1,2	1,3	30,8	1,3	8%
11	12	1,24	32,3	1,24	1,36	29,4	1,36	10%
12	14	1,28	31,3	1,28	1,42	28,2	1,42	11%
13	16	1,32	30,3	1,32	1,48	27,0	1,48	12%
14	18	1,36	29,4	1,36	1,54	26,0	1,54	13%
15	20	1,4	28,6	1,4	1,6	25,0	1,6	14%
16	22	1,44	27,8	1,44	1,66	24,1	1,66	15%
17	24	1,48	27,0	1,48	1,72	23,3	1,72	16%
18	26	1,52	26,3	1,52	1,78	22,5	1,78	17%
19	28	1,56	25,6	1,56	1,84	21,7	1,84	18%
20	30	1,6	25,0	1,6	1,9	21,1	1,9	19%
21	Note: To determine no	t the incremer	ntal effect bety	ween two wei	ghts (E3 vs.B3), but the pp_	ISF effect	
22	for the investigat	ted _weight (I	E3): Enter 0 in	(B3)				
23	Column (H) then	shows the ful	l effect of pplS	SF with the se	lected weight	(E3)		
0.4								

 Now, going with a 50% stronger pp ISF weight of 0.03 (table: E3):

before
$$1,32 = 1 + 0.02 * delta => 0.32 = 0.02 * delta => delta = 16 mg/dl$$

after
$$? = 1 + 0.03 * delta => ? = 1 + 0.03 * 16 = 1.48$$

New effective ISF would be $40 / 1.48 = \frac{27.03}{9}$ mg/dl/U (box F13 in table).

For an intended correction by $-\,10$ mg/dl the <code>insulin</code>Required would now calculate to 10 /

27.03 = 0.370 U, which is 12.1 % more insulin (box H13 in table).

The % effect would be more or also less pronounced, depending how strong bg is rising (**delta**), which is of course the key idea behind pp_ISF.

An **Excel tool** is provided in e-book section 04 (Github/bernie4375) for this calculation. However,

a simple "trial and error" approach as outlined initially in this 4.3.2 section (2 pages before) might be easier to handle.



To summarize, tuning-in the pp_ISF_weight allows the user to define her/his personal sweet spot for ISF aggressiveness in the phase of strongly rising bg.

 • As different kinds of meals will have different **delta** patterns, it can well be that one pp_ISF_weight (as set by you in AAPS preferences) is good for all meals.

634	 Note that a circadian profileISF provides an avenue to still differentiate between e.g.
635	breakfast and lunch response of your autoISF loop.
636 637	4.3.3 Loop states with very little insulin need (iob > iobTH, or 0 %TBR)
638	Normally (except for very low carb meals) the SMBs triggered by bgAccel_ISF_weight and
639	pp_ISF_weight should be sufficient to reach and slightly exceed the iobTH (see <u>section 2.4</u>) so all
640	the other autoISF parameters are relatively unimportant for now.
641	
642	A reason why this can work at all, also for quite a variety of meals, lies in the fact that there
643	is an hourly carb absorption limit of about 30g/h
644	(Reference: Dana
645	Lewis: https://github.com/danamlewis/artificialpancreasbook/blob/master/8tips-and-tricks-
646	for-real-life-with-an-aps.md#heres-the-detailed-explanation-of-what-we-learned. (That limit
647 648	can be lower, e.g. with gastroparesis or certain medications, but that would make things even easier)
649	even easier)
650	So while meals might wildly vary in composition and size: What is digested, and needs insulin in
651	the first ~90 minutes (when FCL tries to catch up with insulin need and differs strongly from HCL,
652	with bgAccel ISF and pp ISF in the leading role), will be relatively closefor meals with similar
653	initial glucose acceleration and rises, anyways
654	
655	The others, low carb with much slower initial acceleration and rise, are easy recognized as
656	different by the loop, see section 4.4 that follows.
657	
658	Depending on the type of meal and "aggressiveness" of your bgAccel_ISF_weight and
659	pp_ISF_weight tuning, the iob will already be so high that, in the phase of decelerated glucose rise
660	towards the peak (the "last part of the rise"), no more insulinRequired is seen by the loop.
661	
662	Therefore the bgBrake_ISF_weight is often unimportant in meals with a relevant carb content.
663	For potential relevance in low carb meals, see section 4.4.
664	
665	
666	4.3.4 "Quality control" on how well your tuning can replace your former HCL bolussing
667	
668	Warning: Occasionally consult the SMB tab to see how your settings really work.
669	A setting (ISF_weight) that is actually set too aggressive might be masked.
670	Tuning only works if the effects of the settings being tuned are not unintentionally limited by
671	other (e.g.,,safety") settings.

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- Also, always look at two or three different meals before deciding whether a tuning "fits" ("good
 - enough" for each of them). You probably will have to iterate back and forth doing this for two or
 - three different kinds of meals ...
 - Case Study 4.1 (Pizza Meal) contains, towards the end, an example how you can go about tuning the weights for various ISF factors of autoISF.
 - Case Study 8.2 shows that it is **not** worth it to seek "optimized" settings based on just one (more extreme) meal.
 - ... until you find one good enough set of settings for all of them. Do not rush this, establishing a solid foundation will be well worth your time.
 - The following sections will deal with similar issues like you were facing in HCL after your given bolus lost much of its power, and SMBs were needed for the "eCarbs".
 - 4.4 Sluggish rise towards a bg peak: bgBrake ISF
 - At a low carb meal, or an attempt at doing a weight reduction diet, (and probably also with gastroparesis, or if you take one of these novel GLP-1 drugs that slow meal absorption -Somebody, please supply a case study!)- the glucose goes up only sluggishly, and iobTH should not be reached at all.
 - In case you exclusively do very slow absorbing meals, you could of course also adjust your iobTH setting low enough to suit your *uniform* situation.
 - Acceleration, and the phase of strong glucose rise, are quickly over at slow-absorbing meals, and there can be:
 - a decelerating bulge of insulin action that projects over an hour or longer. This is where the importance of the **bgBrake ISF** can come in.
 - a bg curve that hovers for an hour or longer around an elevated bg level, because additionally absorbed carbs, and consumption of the moderate SMBs delivered, tend to keep a balance for a while. Dura ISF can deal with this (see next chapter). An example for this is given in Case study 4.2.
 - Note that in some data outputs (e.g. the csv/xls tables coming from the Emulator, e.g. in Case study 4.2, big table at the end there), you will see only "acce_ISF" results.
 - In case of positive acceleration, these are driven by the bgAccel_ISF_weight setting, and results are >1.
 - In case of negative acceleration (decelerating rise), bgBrake_ISF_weight is applied, , and results are < 1. (Example see in graph in <u>section 10.3.3.3</u>).

709 710 In full closed loop, the bgBrake ISF weight is often only about half as large as the 711 bgAccel ISF weight (but that would also depend on your personal diet pattern and 712 eating/digestion speed). Also here, one should approach the tuning gradually, increasing the 713 weight coming from small values. 714 715 Please observe that tuning bgBrake_ISF_weight must strictly be done with types of meals for 716 which there is insulin need at de-celerating but still rising bg. 717 bgBrake ISF is totally irrelevant for hi carb meals where your loop shot over iobTH already 718 by the time your rising towards the bg peak slows down! 719 Likewise, if your initial bgAccel weight is set so strong that your first SMBs catapult you 720 over the iobTH, no matter what type of meal: Then you must first find a reasonable setting 721 for this parameter, one that works "good enough" to control your carb loaded meals, and 722 then see whether there is "room" (and need) for milder loop response at low carb meals. 723 In case you cannot quite get all the ISF_weights "right" so the occasional low carb meal will not get 724 725 over-treated: Avenues to adapt your loop aggressiveness are discussed in section 5. 726 For instance you will be able to (if needed): 727 728 • use a temp. reduced %profile 729 temp. lower iobTH or bgAccel ISF weight 730 construct for yourself a "DIY cockpit" with an extra "snack" or "low carb" button with an 731 underlying suitable Automation 732 733 In the late stage of still rising (!) glucose, the Full Closed Loop typically sharply reduces 734 SMBs already because it is "painfully aware" of the following principal conflict: 735 736 iob (like formerly given in HCL via your bolus) must go high quickly, in order to limit the high 737 However, if there is too much insulin in the system, a hypoglycemia can happen later 738 within the DIA time window, because the loop can, later, only correct to a very limited extent 739 (namely, only to the extent that it can set basal to zero). 740 Therefore, the core problem is that the Full Closed Loop must build up iob very quickly, but 741 not too much, in the initial phase of a meal, and high bg values (out of range, >180 mg/dl) 742 can not always be avoided. 743

4.5 Sluggish rise into a bg plateau, or late plateauing at high bg: 745 dura ISF and bg ISF 746 747 748 749 Depending how your personal diet spectrum looks, you need to tune-in your dura ISF primarily 750 with large hi-FPU meals, and/or for meals at the low carb end of your diet 751 752 4.5.1 dura ISF for sluggish rise into a bg plateau 753 754 A (in that case, often not very high) plateau can form in **low carb meals**, when, basically, carb and 755 insulin "burn rates" might keep a balance over an hour or longer, requiring occasional moderate 756 size SMBs.(See an example in case study 4.2) 757 758 4.5.2 dura ISF for late/high bg plateaus 759 760 With large or high fat/protein meals, often a long high bg plateau is encountered (sometimes 761 associated with 2nd "late, long stretched hill" forming for this, in the bg curve). 762 For such situations, autoISF features the modulation of ISF depending on bg level and on duration 763 of **plateau** formation. 764 765 4.5.3 "One size fits all" -dura ISF 766 767 Absolute "pros" could primarily calibrate their dura ISF for low carb. 768 769 Dura ISF has in-built amplification at higher bg levels. So, effects will automatically be boosted in 770 case much higher plateaus develop after greasy feasts. 771 Should that not per se be sufficient, there is more the DIY "pro" can do: 772 by adding an Automation that gives an extra boost "against" the temporary insulin 773 resistance associated with fats (via increasing the baseline, in terms of a temp.130% profile 774 switch, for instance. Compare at: 775 https://androidaps.readthedocs.io/en/latest/Usage/FullClosedLoop.html#stagnation-at-high-776 bg-values), 777 • or by making additional use of the bg_ISF (or dynamicISF) (-> Tune it in parallel.) 778 The author's preference would be to go via Automation, but only in case the in-built 779 differentiation via bg level make it necessary. 780 781 782

783 4.5.4 Options to limit iob delivered from dura ISF

Rather than relying on your initial tuning to keep you safe from hypos also in the future, there are some extra precautions you could take. Some were discussed in Discord or in dev circles,

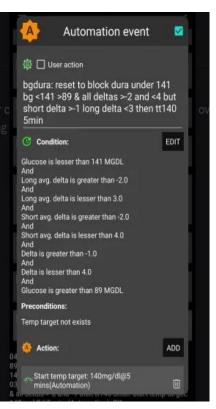
regarding what could be done:

 To limit the danger of going low, it can make sense to design an **Automation** which pauses the delivery of more insulin.

This one was suggested by Alex999

If a glucose plateau built under 140 mg/dl, do not treat via dura_ISF (because the defined Action is to set an elevated TT to a level that will not require more correction insulin.

An alternative Action would be to set, near the actual glucose target, an odd-numbered TT (which blocks any SMB be given, while valid).



- 2) In an autoISF update, the **duration** in which iob is added up could be **capped** after max. 1.5 hours of any "stubborn high".
- 3) Instead of 2), or additionally, the total **iob accrued in that "dura phase" could be capped** by a new related safety setting. It would probably be anchored on iobTH, and could also become a tuneable setting, maybe even a new parameter useable in Automations, too.

4.5.5 How dura ISF works

- 811 Conditions for dura ISF to become active:
- 1) Glucose plateaus, i.e. it is varying within a +/- 5% interval only, and this situation lasted at least for the last 10 minutes => Duration **dura_05** = 10 or more minutes.
 - 2) The average glucose value in this "duration" time window is **avg05** mg/dl. Its elevation relative to bg target determines one of the factors in the equation (eq.3) for dura_ISF, see below.

817 Effect:

The strengthening of ISF is stronger the longer the situation lasts, and the higher the average glucose is above target:

 This can be individually tuned by the duralSF_weight to automatically manage high plateaus in bg values.

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820

How changing the _weight influences the resulting calculated insulinRequired

824 (You can continue 3 pages down at 4.5.6, unless you want to know quantitatively how things work).

825

The developers' documentation (Quick Guide, page 6) https://github.com/ga-

<u>zelle/autoISF/blob/A3.2.0.4 ai3.0.1/autoISF3.0.1 Quick Guide.pdf</u> gives the following equation:

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827

```
dura\_ISF = 1 + \frac{avg05-target\_bg}{target\_bg} * \frac{dura05}{60} * dura\_ISF\_weight  (eq. 3)

profile\_ISF / dura\_ISF = effectively used ISF (sens)
```

(...if the dura influence dominates, and is used as effective ISF. Else, see flowcharts in Quick Guide)

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When looking at the same moment, we can combine the first two factors factors of eq.3 (which vary with bg elevation above bg target, and with how long the plateau already shows) into a term "iF".

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For estimating the effect from using another ISF_weight, we then have two (eq.3), with two unknowns, iF ("cancelling out"), and the sought dura_ISF (that results for the new _weight).

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Example: Your profile ISF is 40 mg/dl/U.

Using dura_ISF_weight of 0.6 you saw effectively used ISF of 30.3 mg/dl/U (box C21 in table on next page) = 40 / 1.32 => factor for dura ISF =1.32.

For an intended correction by -10 mg/dl the insulinRequired would calculate to 10 / 30.3 = 0.330 U.

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842

Now, going with a 33% stronger dura ISF weight of 0.80 (box E3):

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850851

New effective ISF would be 40 / 1.426 = 28.04 mg/dl/U (F21 in table).

For an intended correction by -10 mg/dl the **insulin**Required would now calculate to 10 /

28.04 = 0.357 U, which is 8.1 % more insulin (H21 in table next page).

2 Ins	ofile.ISF (mg/dl/U) sReq (U) @ iF=0 ura_ISF_weight: iF	40 1 0.6			ning -												
3 <u>du</u>	ıra_ISF_weight:	1				dura_ISF tuning -		Ī	4								
-		0.0		see FCL e-Book section 4.5.5		.5.5			4	Α	В	С	D	E	F	G	H
4	ij	0,0			0,8			InsReq e	34	0,966	1,579	25,3	1,579	1,773	22,6	1,773	12,2%
		dura_ISF_old	ISF_old	InsReq_old	ura_ISF_new	ISF_new	InsReq_new	E3 vs.B3	35	0,999	1,599	25,0	1,599	1,799	22,2	1,799	12,5%
5	0	1	40	1	1	40	1	0.0%	36	1,032	1,619	24,7	1,619	1,826	21,9	1,826	12,7%
6	0,033	1,020	39,2	1,020	1,027	39,0	1,027	0,7%	37	1,066	1,639	24,4	1,639	1,853	21,6	1,853	13,0%
7	0,067	1,040	38,5	1,040	1,053	38,0	1,053	1,3%	38	1,099	1,659	24,1	1,659	1,879	21,3	1,879	13,2%
8	0,100	1,060	37,7	1,060	1,080	37,0	1,080	1,9%	39	1,132	1,679	23,8	1,679	1,906	21,0	1,906	13,5%
9	0,133	1,080	37,0	1,080	1,107	36,1	1,107	2,5%	40	1,166	1,699	23,5	1,699	1,932	20,7	1,932	13,7%
10	0,167	1,100	36,4	1,100	1,133	35,3	1,133	3,0%	41	1,199	1,719	23,3	1,719	1,959	20,4	1,959	13,9%
11	0,200	1,120	35,7	1,120	1,160	34,5	1,160	3,6%	42	1,232	1,739	23,0	1,739	1,986	20,1	1,986	14,2%
12	0,233	1,140	35,1	1,140	1,187	33,7	1,187	4,1%	43	1,265	1,759	22,7	1,759	2,012	19,9	2,012	14,4%
13	0,266	1,160	34,5	1,160	1,213	33,0	1,213	4,6%	44	1,30	1,780	22,5	1,780	2,040	19,6	2,040	14,6%
14	0,300	1,180	33,9	1,180	1,240	32.3	1,240	5,1%	45	1,40	1,840	21,7	1,840	2,120	18,9	2,120	15,2%
15	0,333	1,200	33,3	1,200	1,266	31,6	1,266	5,6%	46 47	1,50	1,900	21,1	1,900	2,200	18,2	2,200	15,8%
16	0,366	1,220	32,8	1,220	1,293	30,9	1,293	6.0%	48	1,60 1,70	1,960 2.020	20,4 19.8	1,960 2.020	2,280 2,360	17,5 16,9	2,280 2.360	16,3% 16,8%
17	0,400	1,240	32,3	1,240	1,320	30,3	1,320	6,4%	48	1,70	2,020	19,8	2,020	2,360	16,9	2,360	17,3%
18	0,433	1,260	31,8	1,260	1,346	29,7	1,346	6,9%	50	1,80	2,080	19,2	2,080	2,440	15,4	2,440	17,3%
19	0.466	1.280	31,3	1,280	1,373	29.1	1,373	7.3%	51	2.00	2,140	18,2	2,140	2,520	15,9	2,600	18,2%
20	0,500	1,300	30,8	1,300	1,400	28,6	1,400	7,7%	52	2,10	2,260	17,7	2,260	2,680	14,9	2,680	18,6%
21	0,533	1,320	30,3	1,320	1,426	28,0	1,426	8,1%	53	2.20	2,200	17,2	2,200	2,760	14,5	2,760	19,0%
22	0.566	1.340	29.9	1.340	1,453	27.5	1,453	8.5%	54	2,30	2,320	16,8	2,380	2,700	14,1	2,700	19,3%
23	0.599	1.360	29,4	1.360	1.480	27,0	1.480	8.8%	55	2.40	2,440	16,4	2,440	2,920	13.7	2,920	19,7%
24	0.633	1,380	29.0	1,380	1,506	26.6	1,506	9.2%	56	2,50	2,500	16.0	2,500	3,000	13,3	3,000	20,0%
25	0.666	1,400	28,6	1,400	1,533	26,1	1,533	9.5%	57	3,0	2,800	14,3	2,800	3,400	11,8	3,400	21,4%
26	0.699	1,420	28.2	1,420	1,559	25.6	1,559	9.9%	58	3,5	3,100	12,9	3,100	3,800	10,5	3,800	22,6%
27	0,733	1,440	27,8	1,440	1,586	25,2	1,586	10,2%	59	4,0	3,400	11,8	3,400	4,200	9,5	4,200	23,5%
28	0,766	1,460	27,4	1,460	1,613	24.8	1,613	10,5%	60	4,5	3,700	10,8	3,700	4,600	8,7	4,600	24,3%
29	0,799	1,480	27,0	1,480	1,639	24,4	1,639	10,8%	61	5,0	4,000	10,0	4,000	5,000	8,0	5,000	25,0%
30	0,833	1,500	26,7	1,500	1,666	24,0	1,666	11,1%	62	Note: To determine i	not the increm	nental effect	between two	weights (E3	vs.B3), bu	t the dura_IS	F effect
31	0,866	1,519	26,3	1,519	1,693	23,6	1,693	11,4%	63	for the investig							
32	0,899	1,539	26,0	1,539	1,719	23,3	1,719	11,7%	64	Column (H) the							
33	0.932	1.559	25.6	1.559	1,746	22.9	1,746	12.0%		Use table (I21 x S39					for the obs	served dura	
	,		,	,	,	,	,	,	66	and avg05 =:	> which "iF lin	e" applies ii	n table (A3 x	H61)			

The effects could be more or also less pronounced because the iF factor strongly depends (as in eq.3, see also table on next page) on:

how high bg is above target (avg05)...

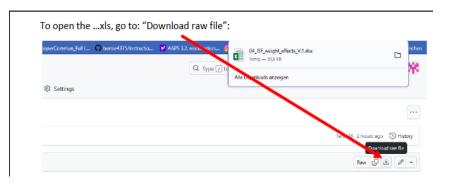
- ...and for how many minutes (dura05).
- Also, do not under-estimate the effect of a low target_bg. Not sure you should, but
 certainly you could add an extra boost from that via an Automation that kicks in a low TT (at
 the "dura situation" that you describe as a triggering condition in your related Automation):

Example: You plateau at 180 mg/dl and your bg target is 100. The first factor in your iF term then is (180-100)/100 = 0.80.

Now, with a TT = 74 mg/dl, that term becomes (180 - 74)/74 = 1.43

(The resulting + 79% in your boost factor does not translate into 79% more insulin - just as we have seen "only" 8% more insulin with a 33% boost in the complete example before).

An **Excel tool** is provided to download as raw filefrom the e-book (Github /bernie4375/section 04) for these complex calculations.



The following table (from this tool) shows examples for combinations of bg target (yellow input field at box J1 resp. box J22), bg plateau level (mg/dl, blue figures), and duration (minutes >10, in violet), the iF of 0.533 in our example above could originate (fields marked green):

4	I	J	K	L	M	N	0	Р	Q	R	S
1	target_bg	100	=>	iF factor	rs resultir	ng at diffe	rent avg0)5 (mg/dl	bg), and	duration	(minutes)
2	dura05;avg05	120	140	160	180	200	220	240	260	280	300
3	10	0,033	0,067	0,100	0,133	0,167	0,200	0,233	0,267	0,300	0,333
4	15	0,050	0,100	0,150	0,200	0,250	0,300	0,350	0,400	0,450	0,500
6	20	0,067	0,133	0,200	0,267	0,333	0,400	0,467	0,533	0,600	0,667
7	25	0,083	0,167	0,250	0,333	0,417	0,500	0,583	0,667	0,750	0,833
8	30	0,100	0,200	0,300	0,400	0,500	0,600	0,700	0,800	0,900	1,000
9	35	0,117	0,233	0,350	0,467	0,583	0,700	0,817	0,933	1,050	1,167
10	40	0,133	0,267	0,400	0,533	0,667	0,800	0,933	1,067	1,200	1,333
11	45	0,150	0,300	0,450	0,600	0,750	0,900	1,050	1,200	1,350	1,500
12	50	0,167	0,333	0,500	0,667	0,833	1,000	1,167	1,333	1,500	1,667
13	55	0,183	0,367	0,550	0,733	0,917	1,100	1,283	1,467	1,650	1,833
14	60	0,200	0,400	0,600	0,800	1,000	1,200	1,400	1,600	1,800	2,000
15	65	0,217	0,433	0,650	0,867	1,083	1,300	1,517	1,733	1,950	2,167
16	70	0,233	0,467	0,700	0,933	1,167	1,400	1,633	1,867	2,100	2,333
18	75	0,250	0,500	0,750	1,000	1,250	1,500	1,750	2,000	2,250	2,500
19	80	0,267	0,533	0,800	1,067	1,333	1,600	1,867	2,133	2,400	2,667
20	85	0,283	0,567	0,850	1,133	1,417	1,700	1,983	2,267	2,550	2,833
21	90	0,300	0,600	0,900	1,200	1,500	1,800	2,100	2,400	2,700	3,000

22	target_bg	74	=>	iF factor	rs resultir	ng at diffe	rent avg0	05 (mg/dl	.bg), and	duration	(minutes)
23	dura05;avg05	120	140	160	180	200	220	240	260	280	300
24	10	0,104	0,149	0,194	0,239	0,284	0,329	0,374	0,419	0,464	0,509
25	15	0,155	0,223	0,291	0,358	0,426	0,493	0,561	0,628	0,696	0,764
26	20	0,207	0,297	0,387	0,477	0,568	0,658	0,748	0,838	0,928	1,018
27	25	0,259	0,372	0,484	0,597	0,709	0,822	0,935	1,047	1,160	1,273
28	30	0,311	0,446	0,581	0,716	0,851	0,986	1,122	1,257	1,392	1,527
29	35	0,363	0,520	0,678	0,836	0,993	1,151	1,309	1,466	1,624	1,782
30	40	0,414	0,595	0,775	0,955	1,135	1,315	1,495	1,676	1,856	2,036
31	45	0,466	0,669	0,872	1,074	1,277	1,480	1,682	1,885	2,088	2,291
32	50	0,518	0,743	0,968	1,194	1,419	1,644	1,869	2,095	2,320	2,545
	55	0,570	0,818	1,065	1,313	1,561	1,809	2,056	2,304	2,552	2,800
33	60	0,622	0,892	1,162	1,432	1,703	1,973	2,243	2,514	2,784	3,054
34	65	0,673	0,966	1,259	1,552	1,845	2,137	2,430	2,723	3,016	3,309
35	70	0,725	1,041	1,356	1,671	1,986	2,302	2,617	2,932	3,248	3,563
36	75	0,777	1,115	1,453	1,791	2,128	2,466	2,804	3,142	3,480	3,818
37	80	0,829	1,189	1,550	1,910	2,270	2,631	2,991	3,351	3,712	4,072
38	85	0,881	1,264	1,646	2,029	2,412	2,795	3,178	3,561	3,944	4,327
39	90	0,932	1,338	1,743	2,149	2,554	2,959	3,365	3,770	4,176	4,581

Still. a "trial and error" approach might be easier to handle (see the following section, 4.5.6):

Off topic: dura_ISF is also very useful in Hybrid Closed Loop. It can be used to elegantly manage, fully automatically, a temporary insulin resistance from fatty acids. Please refer to other papers for details (for instance, sections on FPU and persistent high bg in "Meal Mgt.3...,pdf", available here: https://github.com/bernie4375/HCL-Meal-Mgt.-ISF-and-IC-settings).

4.5.6 Set your dura ISF

Set a **start value of 0.2** for your dura_ISF_weight, and increase only cautiously with an eye on hypo prevention 2-3 hours later.

Caution: Fine tuning this parameter only makes sense *after* you tuned your bgAccel_ISF and pp_ISF well (so your thin yellow insulin activity curve shifts as *far to the left*, towards meal start, as *possible*, which will lower bg peaks and ease the job for dura_ISF).

899	4.5.7 Set your bg_ISF
900	
901	Since in Full Closed Loop we make our loop give us the maximum SMB size it can, at the
902	beginning of a rise, it is crucial to resist the temptation to continue with a particularly strong ISF
903	in the meal phase with the highest glucose values .
904	This is a reason why in Full Closed Loop we do not make much use of the bg_ISF component of
905	autoISF.
906	 Wanting to get most of our insulin from SMBs delivered at fairly low (but beginning-to-rise)
907	bg implies that we do not make ISF weaker at low bg. Under preferences/OpenAPS
908	SMB/autoISF/bg_ISF settings you could set lower ISF_range_weight = 0.0.
909	If you want to analyze in your data, whether you might benefit from a milder ISF at low bg
910	values (e.g. if you often go below target after correction of only mildly elevated bg in the
911	preceding hours), you may want to try lower ISF_range_weight = 0.1 or 0.2. Study the
912	effects from bgISF, and increase, or decrease, the bgISF_weight to fine tune the sought-
913	after affect.
914	• The higher_ISF_range_weight is used when bg is above target, It then strengthens ISF
915	the more the higher the set weight is. 0 disables this contribution, i.e. ISF is constant in the
916	whole range above target.
917	In FCL, this factor should be fairly irrelevant: Near glucose peak, zero-temping usually
918	prevails anyway, so the settings we try might often not be used really by the loop. Very
919	likely, you can live with setting the weight to = 0.0 here, too.
920	If you want to analyze in your data, whether you might benefit from a stronger ISF at high
921	bg values (e.g. if you often remain above target after correction of elevated bg in the
922	preceding hours), you may want to try higher ISF_range_weight = 0.1 or 0.2. Study the
923	effects from bg_ISF, and increase, or decrease, the higher_ISF_range_weight to fine tune
924	the sought-after affect.
925	In case bg_ISF shall play a bigger role in your loop, please consult the related developers'
926	documentation (Quick Guide, page 4) at: https://github.com/ga-
927	zelle/autoISF/blob/A3.2.0.4 ai3.0.1/autoISF3.0.1 Quick Guide.pdf .
928	
929	4.5.8 "Quality control" on your tuning for the later half of your meal time
930	
931	The later stages of meal management (both, in HCL and in FCL) struggle with the problem that
932	there is a hypo danger from the "tail" of insulin activity from earlier SMBs that were needed to

 $fight\ high\ bg\ or\ plateaus\ associated\ with\ temporary\ insulin\ resistance.$

Once your bg sits high, neither you, nor a hybrid closed loop with all the carb info, nor your FCL
can work wonders.
Very important:
 Iterate between 2 or 3 kinds of meals (from your typical spectrum) to find one set of settings that works good-enough for all. That should be possible.
 If you can't make it work for certain meal types, see <u>sections 4.7</u> and <u>5.</u> what you can do then.
Observe hypo trends after meals, and
 resist the temptation to elevate the dura_ISF_weight very high.
• try to stay away from bg_ISF or dynamicISF in Full Closed Loop:
 In FCL you probably can afford to shut bg_ISF entirely off via setting both related _weights to 0.0.
 At least be careful, use small ISF_range_weights and check whether you are happy with the contributions to the effectively used ISFs
 Off topic: If, coming from dynamicISF usage, you stay in Hybrid Closed Loop, but now with autoISF, you probably can use the bg_ISF parameter with higher _weights to emulate what you like to replicate from your dynamicISF experience.
bg highs will take time to resolve.
Interestingly, an after-dinner walk can work wonders sometimes (take glucose tablets along).
Zero-temping and too tightly set safety limits can be stumbling blocks in your tuning project:
 Investigating effects of set ISF_weights is not really possible in periods of zero-temping.
 Too tightly set safety limits "allow" tuning that really is way too aggressive, but gets "cut" by your too-cautious safety settings (e.g. for SMB_range_extention, or for autoISFmax). Aggressive settings then could not come into play most of the time. However, some <i>other</i> time they might come into play, and <i>then</i> produce a hypo 1-2 hours later.
Therefore, carefully study the SMB tab (or better yet, do an emulator based analysis, see
 sections 10-11) to see what the selected weights would do, if there was no zero-temping at the time, and

- whether you bump into a set limit already (if your bgAccel_ISF_weight makes you exceed
 allowed max. SMB size, then further tuning your settings only makes sense with either
 allowing bigger SMBs, or limiting bgAccel_ISF_weight to a lower number at which you will
 not frequently bounce into the SMB limit)
 - at which **other** times (rather than the one you currently look at and try to improve) that selected setting might backfire

973

976 4.5.9 How your "UAM" loop concludes insulin need for your un-declared carbs 977 978 The UAM Full Closed Loop doesn't get any information from you as to how many grams of carbs 979 will be there, for absorption. 980 981 Looking back 982 983 For the recent 5-minute segments, the UAM oref(1) loops can precisely calculate how many grams 984 of carb "must have been digested" based on the CGM values seen, and your IC and ISF profile 985 parameters 986 For more detail see chapter 1.2 on how dynamic carb absorption is calculated, in "Carb ratio (IC, 987 CIR)...pdf" at: https://github.com/bernie4375/HCL-Meal-Mgt.-ISF-and-IC-settings. 988 989 Looking into the next minutes, hour 990 991 However, here we worry about the late meal stage, and our FCL has gotten no information from us 992 about how many grams in total were eaten, and certainly we do not bother to give eCarbs with 993 estimated **absorption times** (that are so essential in iOS Loop). 994 995 So, in FCL you leave your loop without knowledge when your steady-state max carb absorption 996 phase... 997 o the earlier mentioned 30g/h, or 998 o with gastroparesis, or if on GLP-1 drug treatment, probably on a lower g/h level sometimes prolonged ("faked") by a brief episode of insulin resistance to fats 999 1000 ...might end. Nor, whether extra carbs were added, later, or "FPUs are lurking". 1001 1002 The FCL now needs to provide desired amounts of insulin, while facing a potentially induced hypo 1003 danger later (considering the DIA of all the insulin still active). 1004 1005 Fortunately, the UAM Full Closed Loop is *not completely clueless* regarding how carb absorption 1006 will continue: 1007 1008 It will work with a **prediction** of *further* carb absorption, building on the **carb deviation**s 1009 (=calculation of how much got absorbed in the past 5 minute segments), and phase out further 1010 *expected* carb decay in the course of the next 1 to max 3 hours. 1011 1012

1013	For more detail see
1014	• https://openaps.readthedocs.io/en/latest/docs/While%20You%20Wait%20For%20Gear/Und
1015	erstand-determine-basal.html#understanding-the-basic-logic-written-version
1016	• or do a real-time study with (screenshots from) your SMB tab info.
1017	
1018	Discussion
1019	
1020	This UAM prediction about further carb absorption can be worse, but can also be better than a
10211022	prediction based on the user's "e-Carb" input as done in Hybrid Closed Loop.
1023	In any case, and even when having perfect knowledge about how exactly the carbs fade out in the
1024	next hours, there would still be a principal problem for the loop: Heavy insulin "fire" against highs
1025	will not work immediately (depending on the insulin's time-to-peak), and notably it comes with a
1026	significant hypo danger from the "tail" of insulin activity.
1027	A big bolus, or also a series of boli, will rarely work exactly for several hours matching the
1028	absorption of carbs (from what, how much and how fast the user ate).
1029	
1030	Off topic closing remark: With meticulous attention to all carb-related profile parameters, and daily
1031	inputs on amounts and absorption times, plus some "mindfulness" as to which diet habits disturb the
1032	possible balance, there are "pro" loopers (notably on iOS Loop) who achieve impressive %TIR
1033	(and % in tight range) performance. – The author consciously chose the other way, to put substantial
1034	effort into a personalized upfront system calibration, and work with a oref(1) algo system that allows
10351036	(nearly) every day hands-off FCL.
1030	4.6 Tuning your initial settings
1038	
1039	Be pro-active: The earlier large SMBs come (driven by bgAccel ISF and pp ISF)
1040	Note: Also your CGM smoothing may play a role here, that you may want to look into!
1041	the less high the overall increase in BG will be, and (provided you set a proper iobTH)
1042	the lesser the risk will be for a hypo after the meal.
1043	
1044	Therefore, put most of your FCL tuning effort into determining suitable weights for
1045	bgAccel_ and for pp_ISF, and for finding a suitable iobTH_percent.
1046	
1047	Low carbers probably should pay more attention on dura_ISF, besides seeing to it that
1048	bgAccel_ISF is not too aggressive (see above, section 4.2.5 and case study 4.2).
1049	

1050	Later, your "FCL cockpit" will give you access to temporarily modulate these essential
1051	parameters (see section 5.2.), providing you an opportunity
1052	 in your tuning phase, for more research on the fly, so to speak
1053	 everyday, for temp. adaptations to altered insulin sensitivity, or to special
1054	disturbances (should you, occasionally, see a need).
1055	
1056	After you tuned your initial settings well, there should rarely arise a need for "fine tuning" later,
1057	see section 8 and case study 8.2!
1058	See Section 6 and edge study 6.2.
1059	The experience of the author is that it is possible to tune the above mentioned weights for very
1060	different meals in such a way that the glucose almost always remains acceptably in range.
1061	
1062	However, if you come to the conclusion that differentiated settings (for different meals or meal
1063	time clusters) would be easier to establish, and/or work better for you, the following sections
1064	suggest many options you could try and use.
1065	
1066	4.7 Maneuvering through more complex scenarios
1067	
1068	You now can move on, to accommodate more complex scenarios.
1069	
1070	4.7.1 Complex meal spectrum
1071	
1072	• Especially if you are a bit shy of using the Emulator (section 10 and 11) for really detailed
1073	analysis, it can well be that you will not hit one real good system calibration (section 4) for your
1074	entire range of diets (e.g. your meal spectrum at all your lunches).
1075	 Note that between meal time slots (e.g. breakfasts vs lunches) you should differentiate
1076	via different "circadian" <i>profile_ISF</i> s (with which the autoISF-effects always multiply).
1077	 So, in case you occasionally run out of range (bg =70180 mg/dl), your options to prevent,
1077	react, or improve are:
1079	 accepting a few % higher time outside of range for that day (and, if feasible, in the
1080	future avoiding what seemed to have caused it)
1081	o taking a snack (whenever you tend to go low from the "tails" of insulin activity that was
1082	required to fight a peak)

108310841085	 doing a manual "tweak" (if you can think of one in time), to manage the problem manually. For example, briefly going into an odd TT (=temp. blocking more SMBs) can be a very easy-to-handle remedy, sometimes
1086 1087 1088 1089	 define a User Action Automation, and provide an extra "cockpit button" to announce a meal <i>outside of your usual spectrum</i>, so it will automatically be treated differently by your FCL (as you defined in your Automation; example: <u>Case study 5.2</u>). temporarily resorting to "your old" hybrid closed loop.
1090 1091 1092 1093	Instead of accepting such instances, you could launch "improvement projects", that refine your initial tuning (section 4. and sections 8 and 9)
1094 1095	Note, though, that it could be near-impossible to fine-tune <i>if your basics never were "right"</i> and you got lost in a maze of errors and counter-errors.
1096	In that case, only a fresh start might convincingly help.
1097 1098 1099	4.7.2 Complex tasks aside from managing meals
1100 1101 1102 1103	To deal with <i>different</i> disturbances than presented by your meal spectrum (that you were calibrating for in this section 4), there will be other instances where temporary modulations of your FCL will be needed.
1104 1105	You have a variety of options to deal with that, and this will be the topic in <u>section 5.</u>
1106 1107 1108 1109 1110 1111 1112	It is suggested to do major exercise still <i>in your hybrid closed loop</i> setting, <i>until</i> you have your FCL up and running for meals on normal days with no or only moderate exercise. Later, implement extras as discussed in <u>section 6</u> to fully implement your FCL.

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For general loop settings for FCL with autoISF, please consult $\underline{\text{sections } 2.1 - 2.6}$

The following table gives a comprehensive approach how to tune your autoISF FCL better, when you see any of the problems as in *column A*.

Make sure to work through the problems in the order given in this table.

4	Α	В	С	D		
1	Notes:1. if not entering	carbs the	n "several hours after eating" can be applied in	stead of "with zero COB		
	The problem	Y/N	Possible solution	Based on what logic		
Ť	,,					
ı	Are you having perods with zero COB and negative IOB?	Yes	Decrease basal a bit over these periods, wait a couple of days to evaluate and return here	Oref is backing off your basal to try and hold your BG steady so ba is probably too high		
•	and negative IOB?		Continue with this sheet			
	Are you having perods with zero COB	Yes	Increase basal a bit over these periods, wait a couple of days to evaluate and return here	Oref is increasing your basal to try and hold your BG steady so		
	and positive IOB?		Continue with this sheet	is probably too low		
[Do you get roller coaster BGs?	Yes	Increase profile ISF a bit (make weaker). Also, check your set iobTH%. Reevaluate over a couple of days, return here	BG goes over target -> with too-strong ISF, too much insulin is or resulting in BG below target. Long zero-temping required, but resulting the BG next		
		No	Continue with this sheet			
H	After your BG went high, do you get a hypo later (from	Yes	Reduce autoISF "weights" a bit, reevaluate over a couple of days, return here	ISF is reduced (strengthened) too much by autoISF as you move a from target. Reducing the proper ISF_weight will reduce that impact Use the autoISF history table for more information about which is causing the issue: Preferably, "shave of late", at DURA or BG;		
	remaining insulin 'tail" at zero COB)?	No	Continue with this sheet	or ACCEL weights might need reducing, too.		
When you start eating with BG around target and iob near zero, do you get a hypo (in the first 2 hours)?		Yes	Check whether you can reduce your set iobTH%, or SMB range extention, or autoISFmax. Decrease PP (and maybe also ACCEL) "weight", and increase profile carb ratio a bit (make it weaker, so you see the insulin needed relate well to the g of carbs digested by the time you see a hypo tendency). Reevaluate over a couple of days, return here	You're getting too much insulin to cope with the bg rises from the cyou've eaten. Taming the loop can be done in several ways: Weakening profile carb ratio and ISF would work in all situations. Lowering ACCEL and/or PP autoISF "weights" would exactly weak loop over-response inearly stage of BG rise. And the 3 first-mentioned safety settings just limit excesses.		
ļ		No	Continue with this sheet			
When you eat, is iob building up too slow, with a lot of it added at bg peak or even still when BG reduces?		Yes	Check whether you must increase your set iobTH%, or SMB range extention, or autoISFmax. Increase PP "weight", and decrease profile carb ratio a bit (make it stronger, so you see the insulin needed relate well to the g of carbs digested). Reevaluate over a couple of days, return here	You're not getting enough insulin to cope with the bg rises from the carbs you've eaten. If your loop bounces into any of the 3 first-mentioned limits, "tuning more aggressive" might simply not work. Making the loop more aggressive can be done in several ways: Strengthening (lowering) profile carb ratio and ISF would work in all situations. Elevating autoISF "weights" (notably PP) could exactly strengthen letesponse in instances where needed (where the respective "weight"		
8	No		Continue with this sheet	playing a role).		
9 Does BG stay high Yes for long periods?		Yes	Increase DURA "weight" a bit. Reevaluate over a couple of days, return here	DURA reduces ISF (makes it stronger) if BG stays high for longer periods than it should Reminder: Follow this sheet in order! This "last job", dealing with		
9	ior long periods:	1	L	persistent highs via DURA, is much easier after you sorted out how build-up iob faster, so not to go super high in the first place.		
_	lor long periods:	No	Have a short holiday with lots of carbs	build-up lob faster, so not to go super high in the first place.		
0	Further reading:	No	Have a short holiday with lots of carbs	build-up lob faster, so not to go super high in the first place.		
20				A3.2.0.4 ai3.0.1/autolSF3.0.1 Quick Guide.pdf		