



6. İÇ HASTALIKLARI KONGRESİ

26 - 29 Nisan 2023
Hilton Bakırköy - İstanbul



Sağlık Bilimleri Üniversitesi



İç Hastalıkları Uzmanlık Eğitim
Araştırma Derneği



GLP1 tedavisi insülin alternatifi midir?

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Ankara Güven Hastanesi

Endokrinoloji ve Metabolizma Hastalıkları Bölümü



Çıkar çelişkisi beyanı

- Danışma kurulu
 - Novonordisk, Sanofi, AstraZeneca, Eczacıbaşı, Econix, Roche Diagnostics
- Klinik araştırma yürütücülüğü
 - Novonordisk, Novartis, Sanofi
- Kongre Desteği
 - Bilim ilaç, Novonordisk, AstraZeneca

İçerik..

- «GLP1 tedavisi insülin alternatifi midir?» sorusuna hızlı cevap
- Kanıtlar
- GLP-1 agonisti tedavisinde püf noktalar



GLP-1 Agonisti insülin alternatifi midir?

İnsülin tedavisi endikasyonları

- Tip 1 Diyabet
- Hiperglisemik acil durumlar
- Diyabetli bireyde perioperatif izlem
- Gebelik diyabeti

- Tip 2 Diyabetli bireyde tedaviye yetersiz yanıt
 - Diyet + Egzersiz + Oral antidiyabetikler

HAYIR

EVET

Tip 2 Diyabet güncel tedavi yaklaşımı

Yaşam biçimi yönetimi + Oral antidiyabetik
Metformin (Çoğu defa)

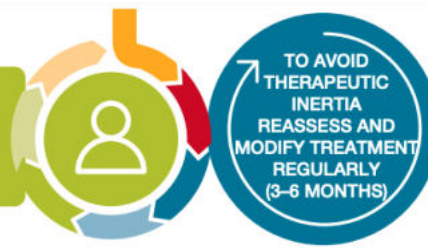


Oral kombinasyon
Endikasyon, Etki, Yan etki, Maliyet



Enjektabl tedavi
GLP-1 reseptör agonisti
İnsülin

Use principles in Figure 9.3, including reinforcement of behavioral interventions (weight management and physical activity) and provision of DSMES, to meet individualized treatment goals



If injectable therapy is needed to reduce A1C¹

Consider GLP-1 RA or GIP/GLP-1 RA in most individuals prior to insulin²
INITIATION: Initiate appropriate starting dose for agent selected (varies within class)
TITRATION: Titrate to maintenance dose (varies within class)

If already on GLP-1 RA or dual GIP and GLP-1 RA or if these are not appropriate OR insulin is preferred

If above A1C target

Add basal insulin³
Choice of basal insulin should be based on person-specific considerations, including cost. Refer to **Table 9.4** for insulin cost information. Consider prescription of glucagon for emergent hypoglycemia.

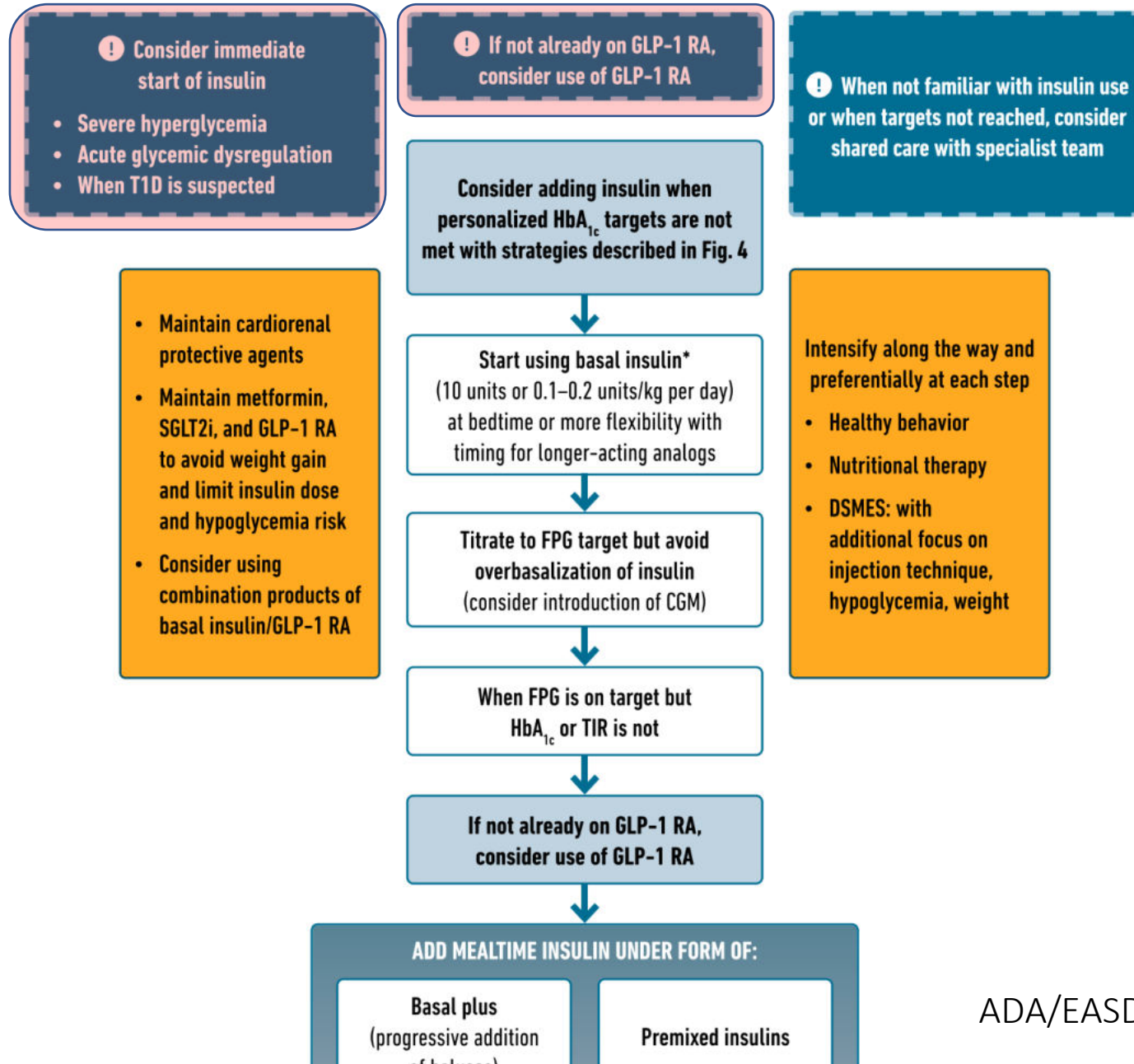
Add basal analog or bedtime NPH insulin⁴

- INITIATION:** Start 10 units per day OR 0.1–0.2 units/kg per day
- TITRATION:**
- Set FPG target (see Section 6, “Glycemic Targets”)
 - Choose evidence-based titration algorithm, e.g., increase 2 units every 3 days to reach FPG target without hypoglycemia
 - For hypoglycemia determine cause, if no clear reason lower dose by 10–20%

Assess adequacy of basal insulin dose

Consider clinical signals to evaluate for overbasalization and need to consider adjunctive therapies (e.g., basal dose more than ~0.5 units/kg/day, elevated bedtime–morning and/or post–preprandial differential, hypoglycemia [aware or unaware], high variability)

PLACE OF INSULIN¹



İnsülin ve GLP-1 agonisti karşılaştırma

Glukoz düşürücü
etki

Metabolik ve Kardiyovasküler

GLP-1 Reseptör Agonistleri

Kısa etkili



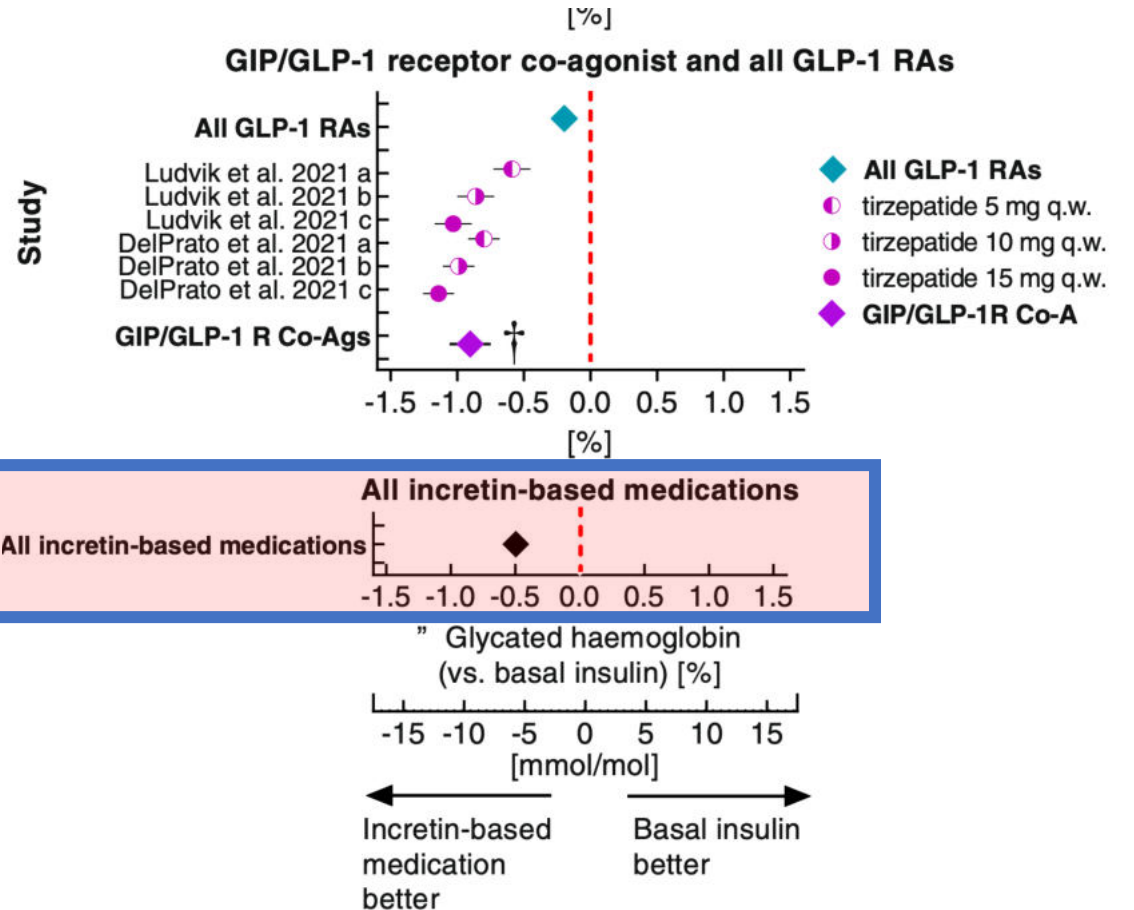
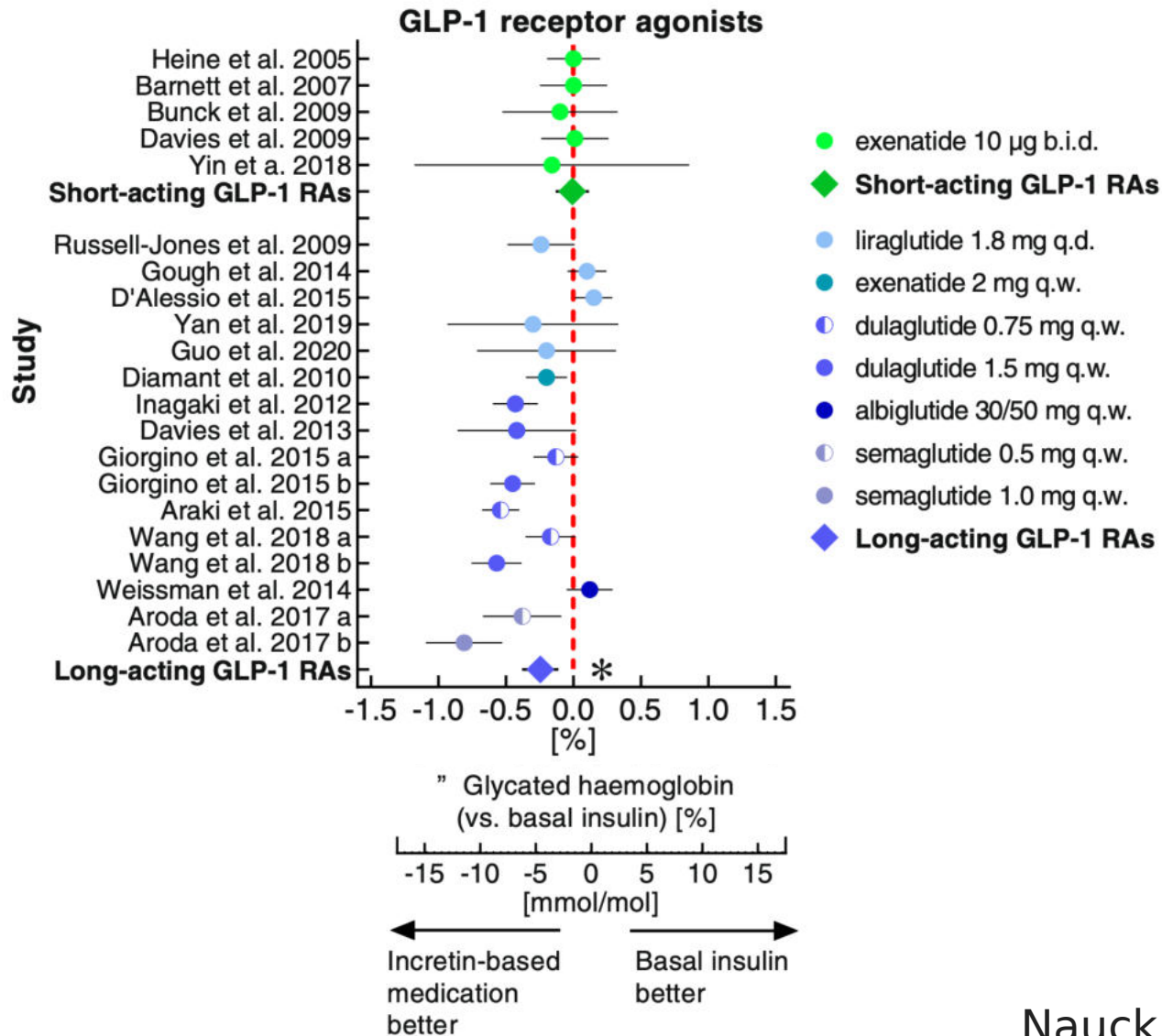
Exenatide
b.i.d.
Byetta®

γ -1/GIP dual agonistler

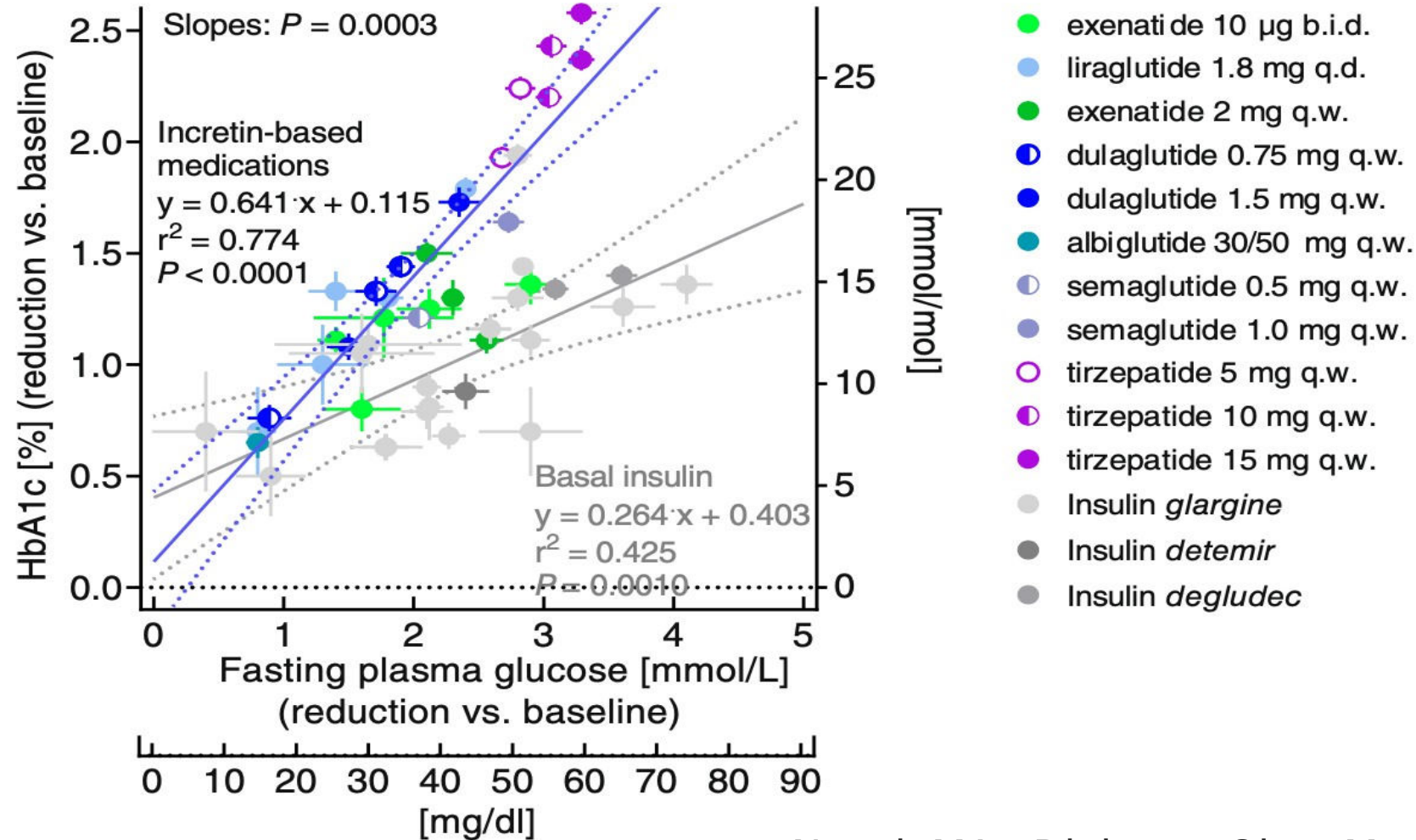


Tirzepatide
Mounjaro

A1c üzerindeki etkinlik



Açlık glukozu üzerindeki etkinlik



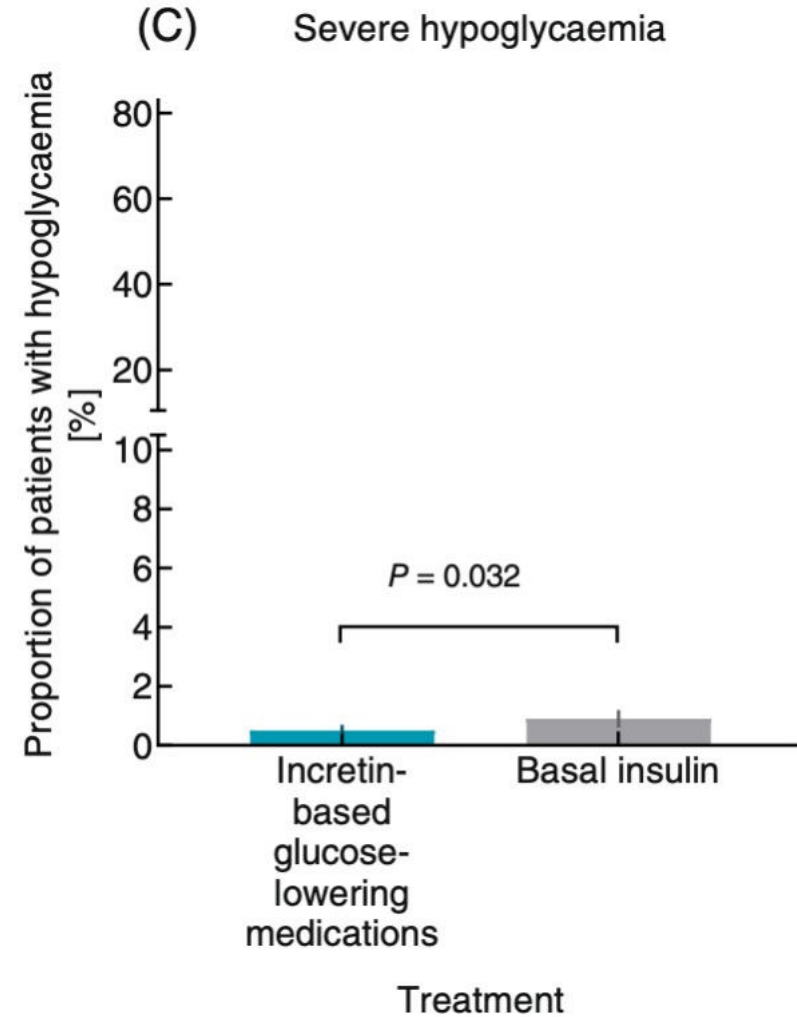
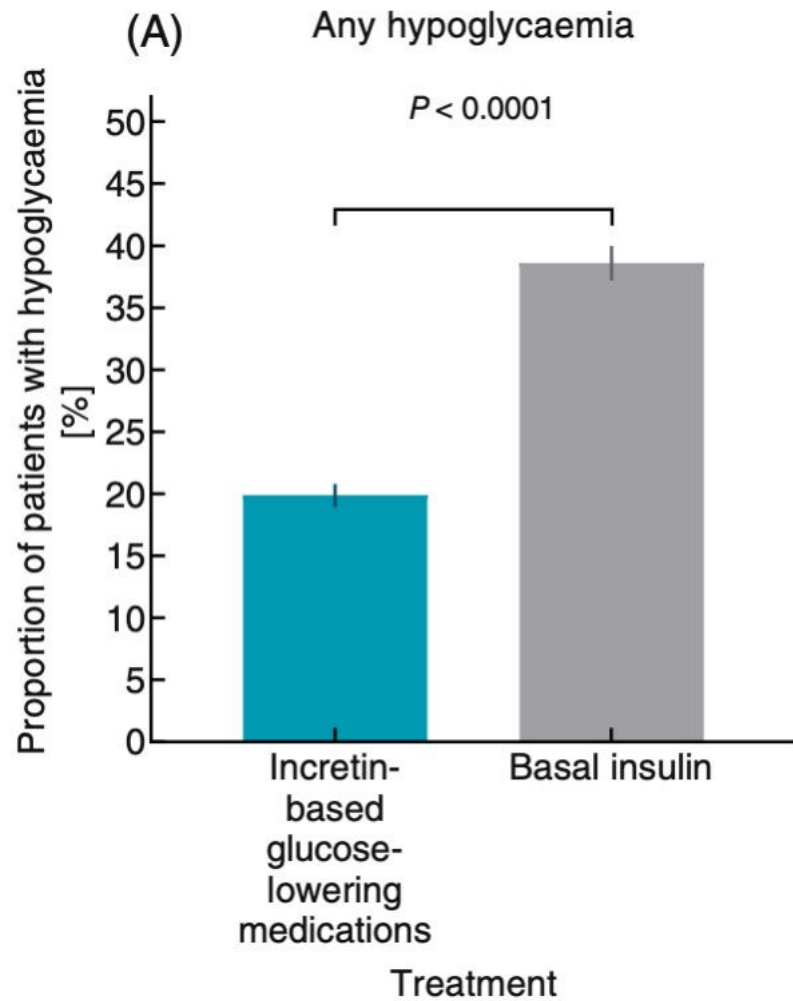
İnsülin ve GLP-1 agonisti karşılaştırma

Yan Etki

Glukoz düşürücü
etki

Metabolik ve Kardiyovasküler

Hipoglisemi riskii..

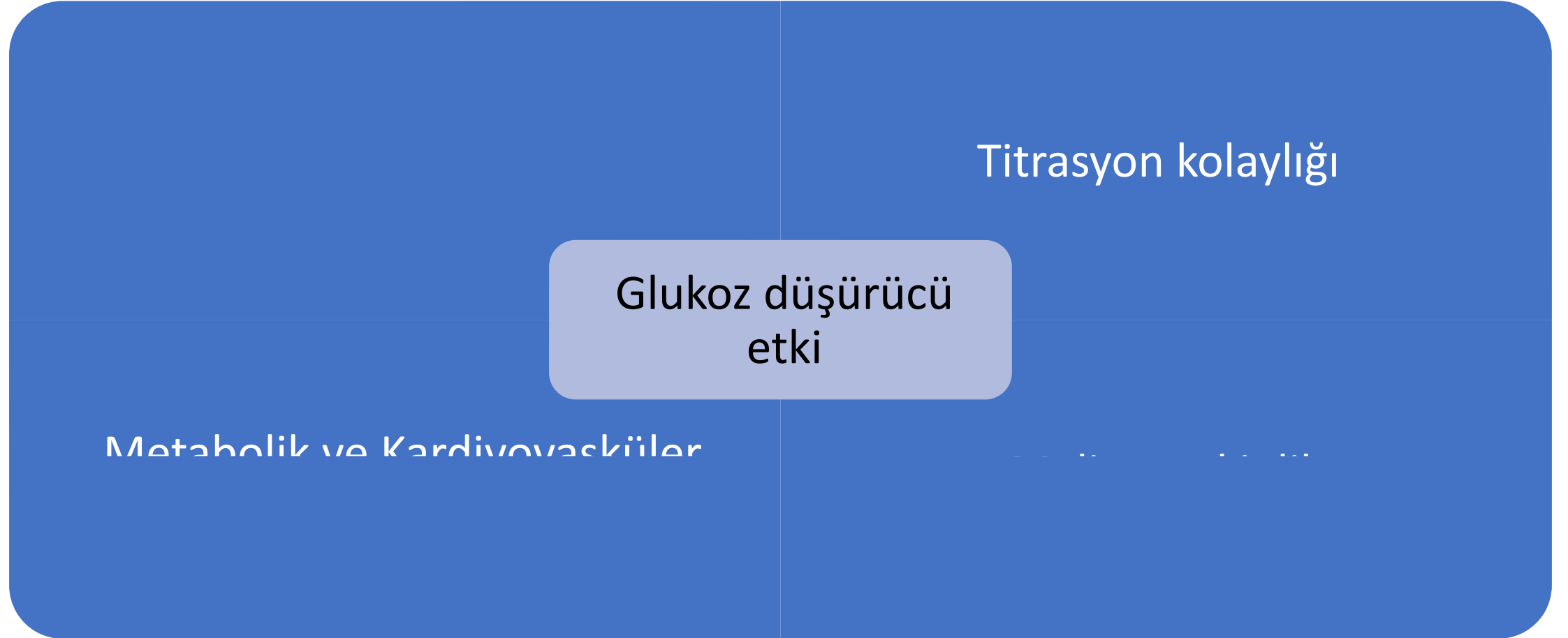


Gastrointestinal yan etkiler..

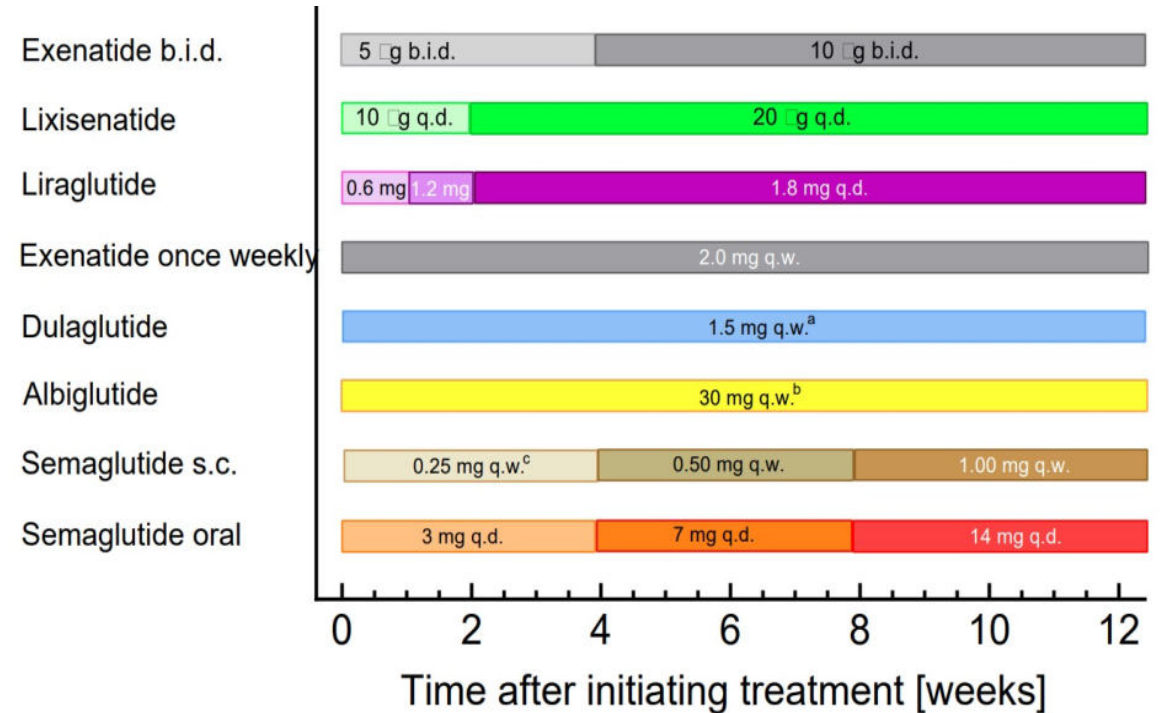
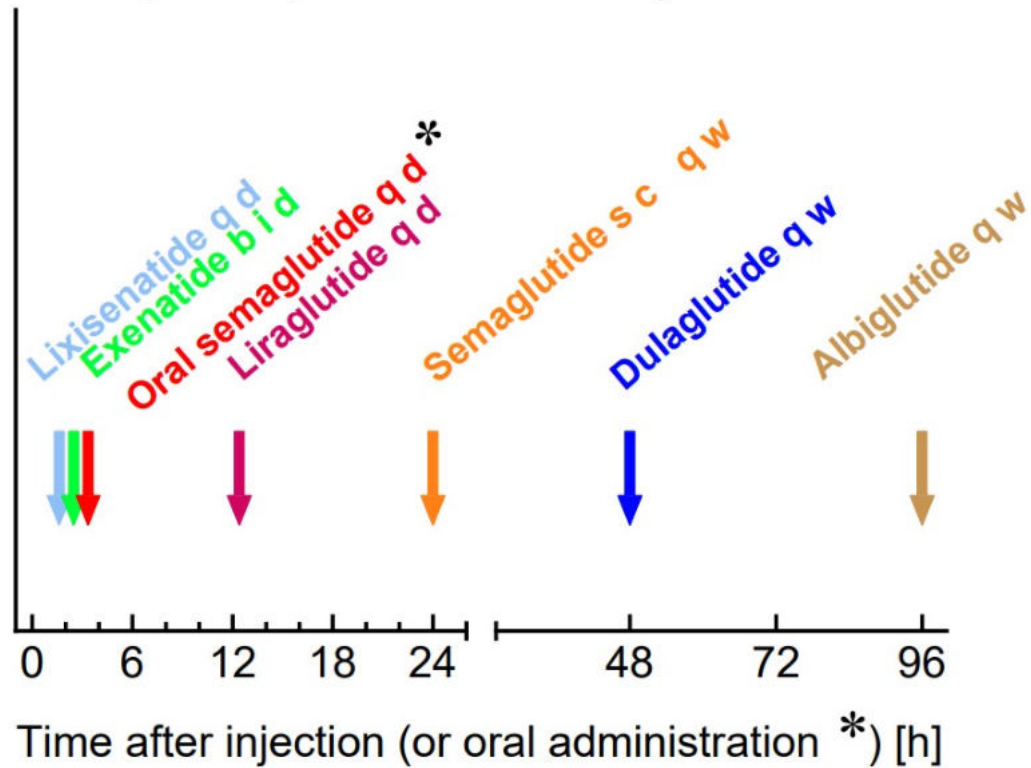
Supplementary Table 3. Comparison of the proportion of patients reporting “gastrointestinal” adverse events (nausea, vomiting, diarrhoea) in head-to-head trials with incretin-based glucose-lowering medications (GLP-1 receptor agonists and the GIP/GLP-1 receptor co-agonist tirzepatide)

| | Number of studies reporting this adverse event | Number of patients reporting this adverse event | Number of patients exposed to the drug(s) of interest | Proportion reporting this adverse event [%] | Number of studies reporting this adverse event | Number of patients reporting this adverse event | Number of patients exposed to the drug(s) of interest | Proportion reporting this adverse event [%] | Number of studies reporting this adverse event | Number of patients reporting this adverse event | Number of patients exposed to the drug(s) of interest | Proportion reporting this adverse event [%] |
|---|--|---|---|---|--|---|---|---|--|---|---|---|
| Adverse event of interest | Nausea | | | | Vomiting | | | | Diarrhoea | | | |
| Medication class(es) analyzed | | | | | | | | | | | | |
| Short-acting GLP-1 RAs | 3 | 289 | 561 | 51.5 | 2 | 61 | 407 | 15.0 | 2 | 46 | 400 | 11.5 |
| Long-acting GLP-1 RAs | 11 | 663 | 4158 | 15.9 | 10 | 255 | 3654 | 7.0 | 9 | 508 | 4190 | 12.1 |
| All GLP-1 RAs | 14 | 925 | 4719 | 20.2 | 12 | 316 | 4061 | 7.8 | 11 | 554 | 4590 | 12.1 |
| Tirzepatide | 2 | 351 | 2072 | 16.9 | 2 | 163 | 2072 | 7.0 | 2 | 351 | 2072 | 16.9 |
| All incretin-based glucose-lowering medications | 16 | 1303 | 6791 | 19.2 | 14 | 479 | 6133 | 7.8 | 13 | 905 | 6662 | 13.6 |
| Comparison (incretin-based glucose-lowering medication vs. basal insulin) | Relative risk | 95 % confidence interval | | Significance (p-value) | Relative risk | 95 % confidence interval | | Significance (p-value) | Relative risk | 95 % confidence interval | | Significance (p-value) |
| Short-acting GLP-1 RAs | 6.3 | 4.4-9.0 | | < 0.0001 | 3.9 | 2.2-6.8 | | < 0.0001 | 1.9 | 1.2-3.1 | | 0.011 |
| Long-acting GLP-1 RAs | 6.2 | 4.9-7.9 | | < 0.0001 | 3.2 | 2.3-4.3 | | < 0.0001 | 2.7 | 2.2-3.3 | | < 0.0001 |
| All GLP-1 RAs | 6.1 | 4.9-7.4 | | < 0.0001 | 3.7 | 2.8-4.8 | | < 0.0001 | 2.6 | 2.1-3.1 | | < 0.0001 |
| Tirzepatide | 6.1 | 4.8-10.1 | | < 0.0001 | 4.4 | 3.0-6.5 | | < 0.0001 | 3.6 | 2.7-4.7 | | < 0.0001 |
| All incretin-based glucose-lowering medications | 6.2 | 5.3-7.1 | | < 0.0001 | 3.9 | 3.1-4.8 | | < 0.0001 | 2.9 | 2.3-3.3 | | < 0.0001 |

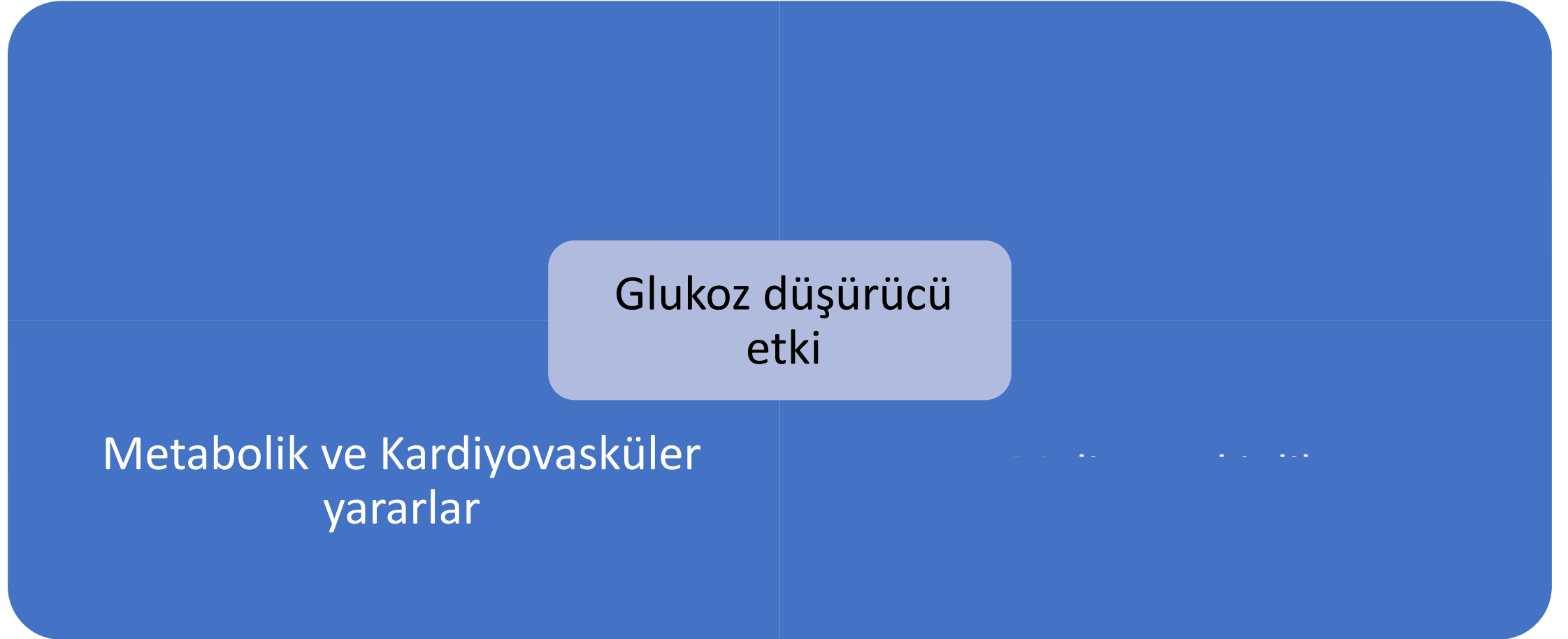
İnsülin ve GLP-1 agonisti karşılaştırma



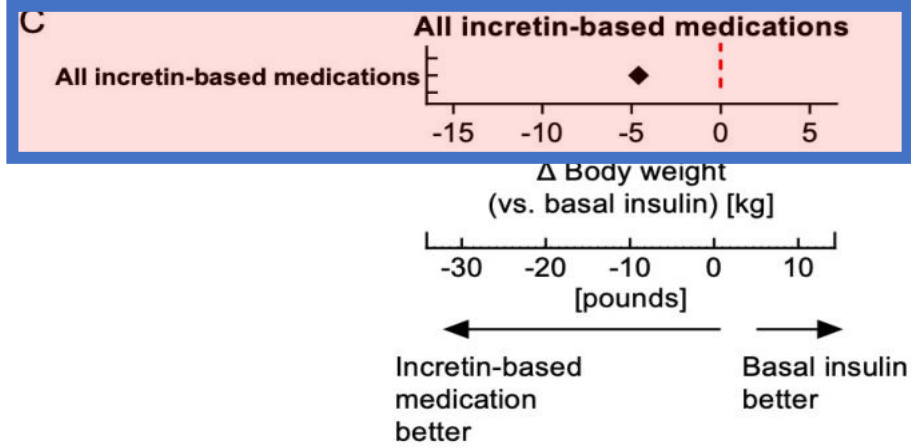
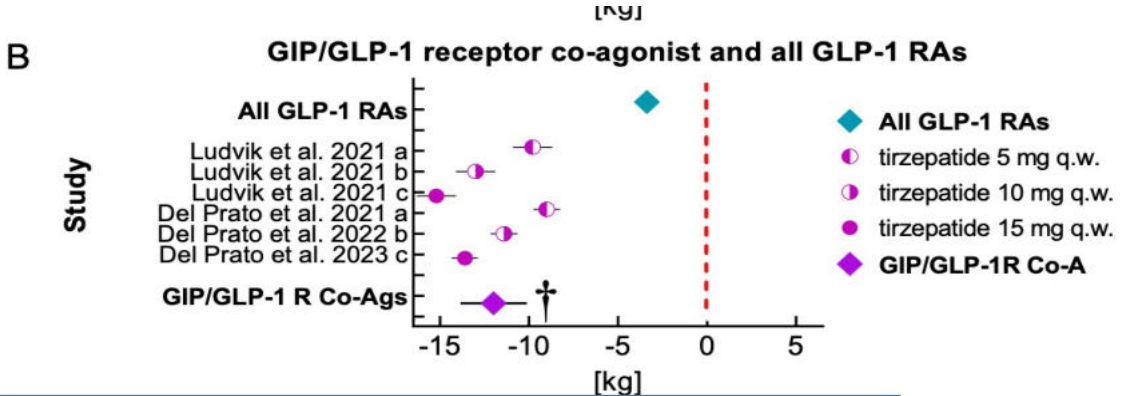
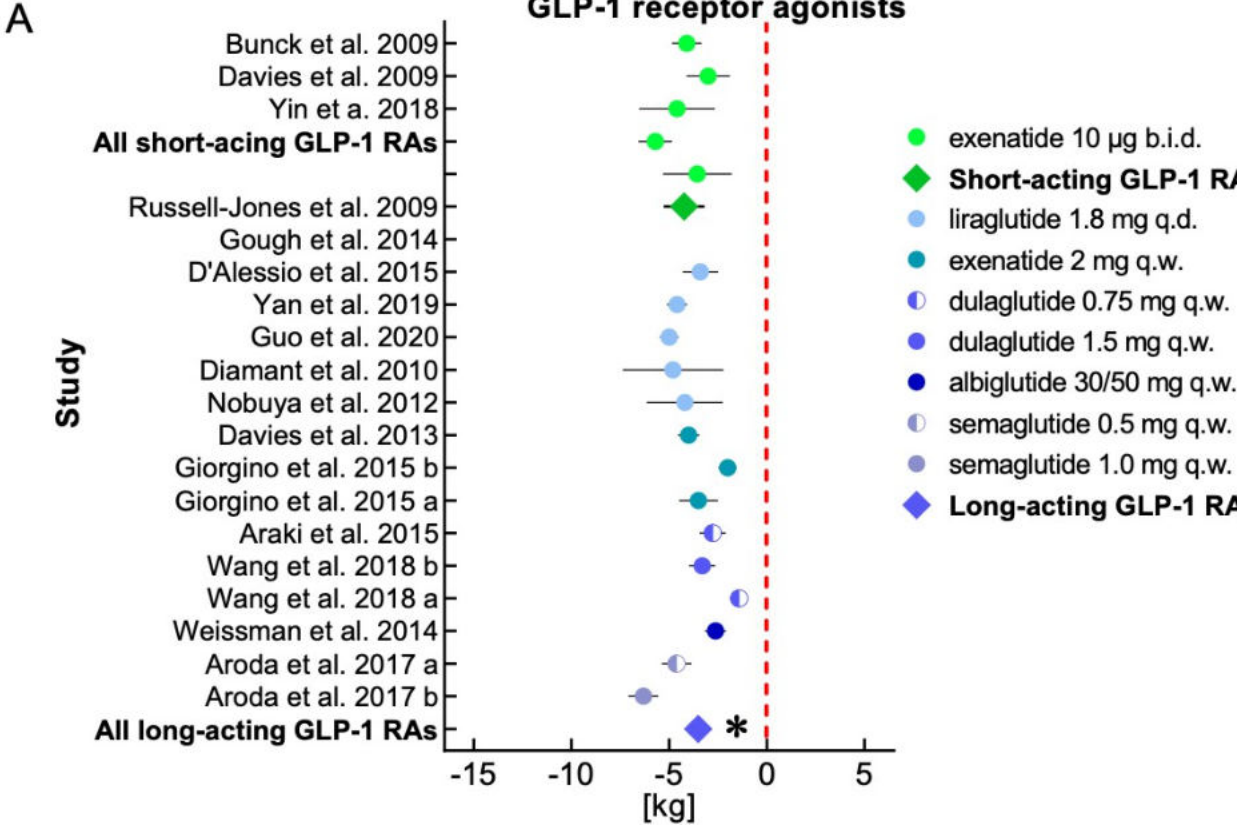
GLP-1 titrasyonla ilgili ip uçları..



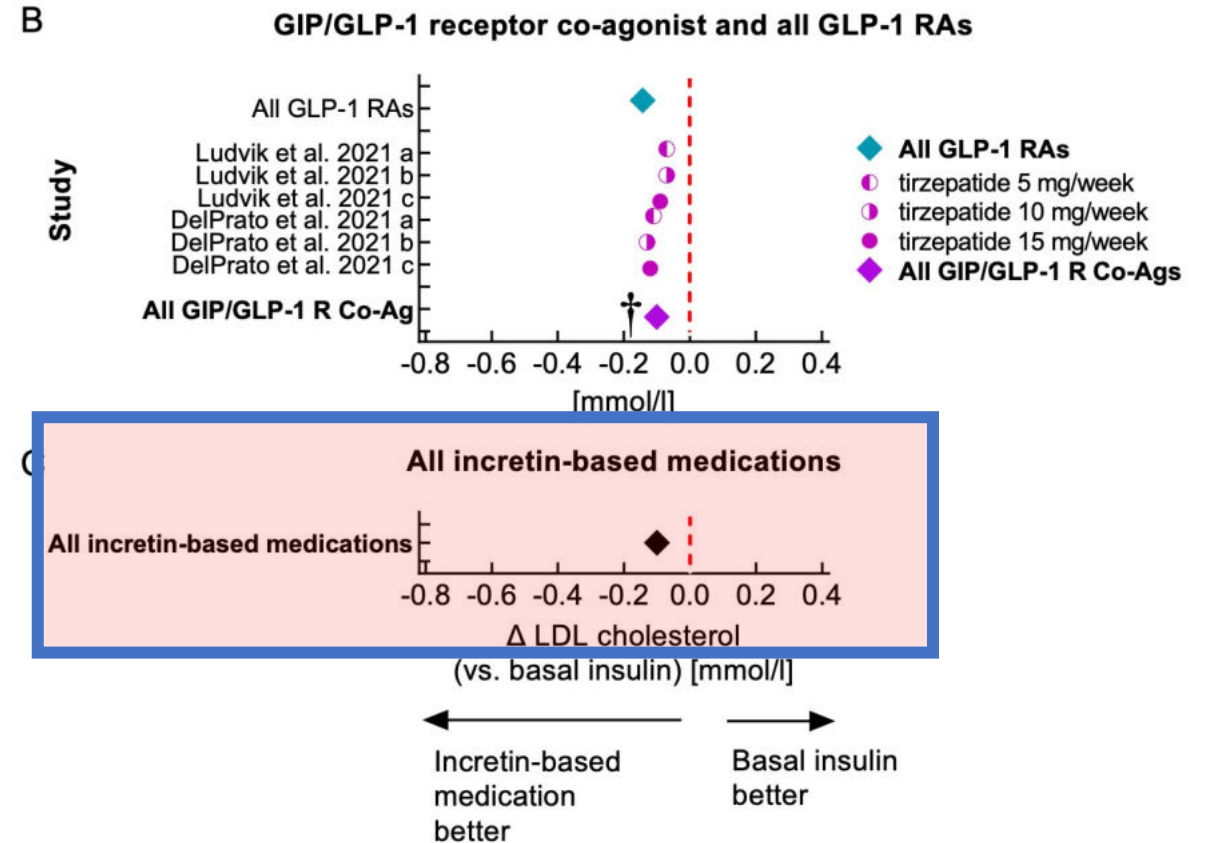
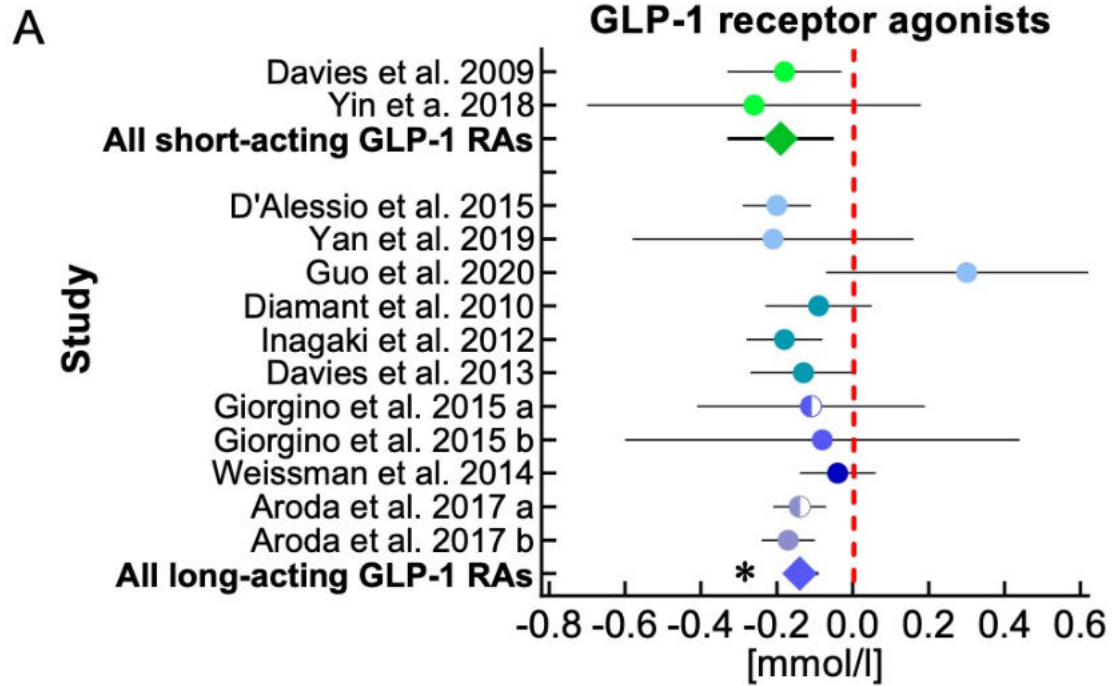
İnsülin ve GLP-1 agonisti karşılaştırma



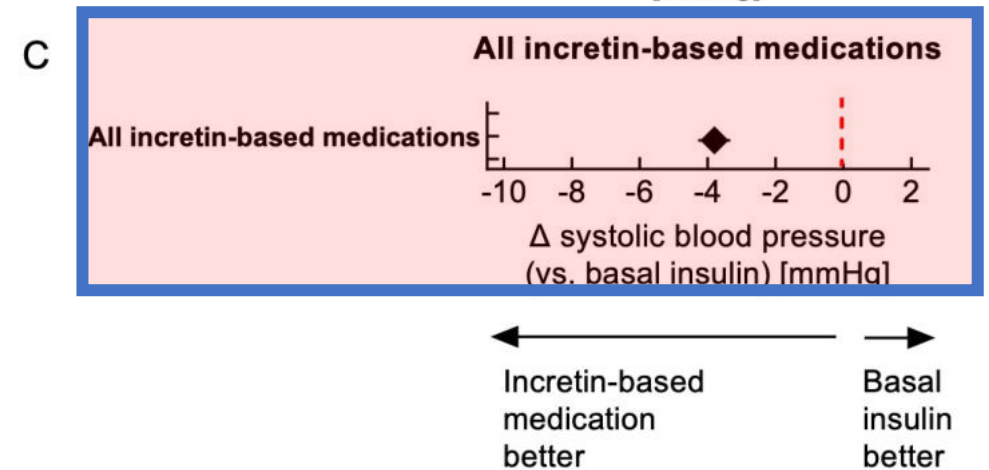
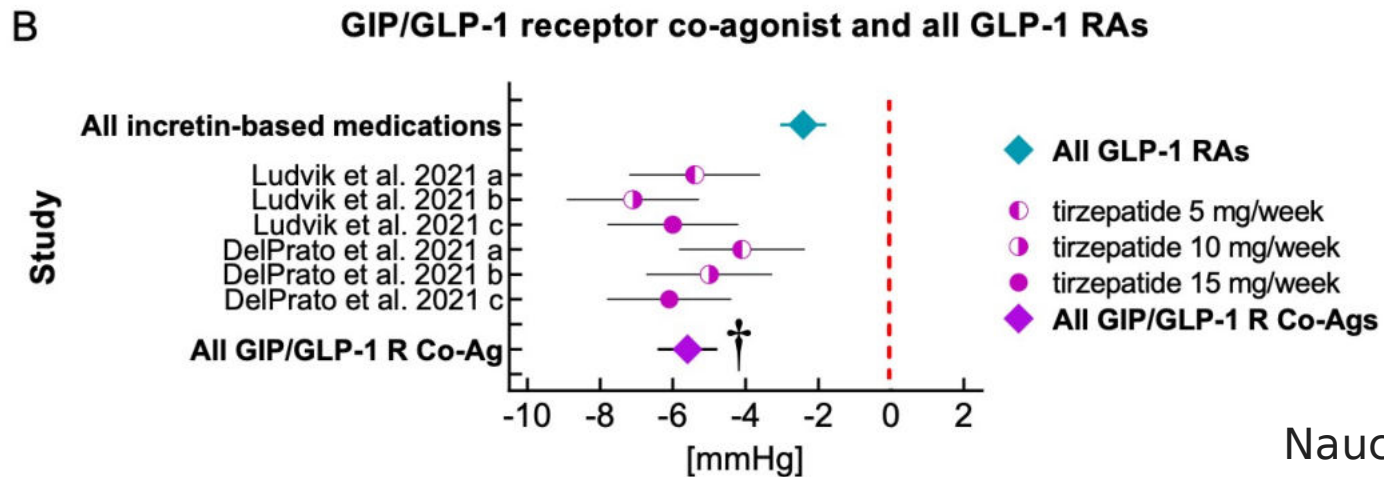
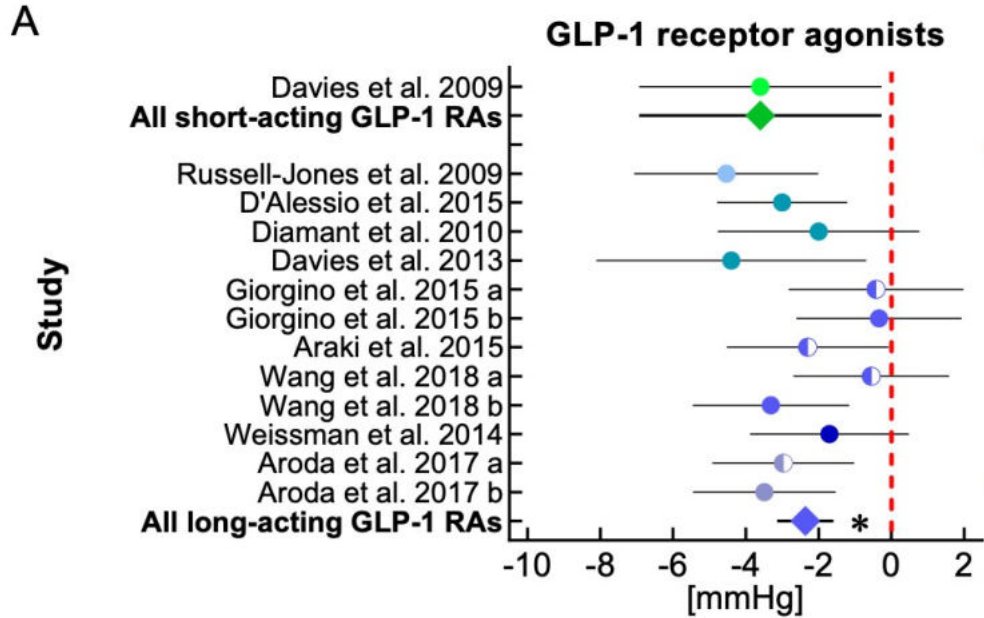
Kilo kontrolü



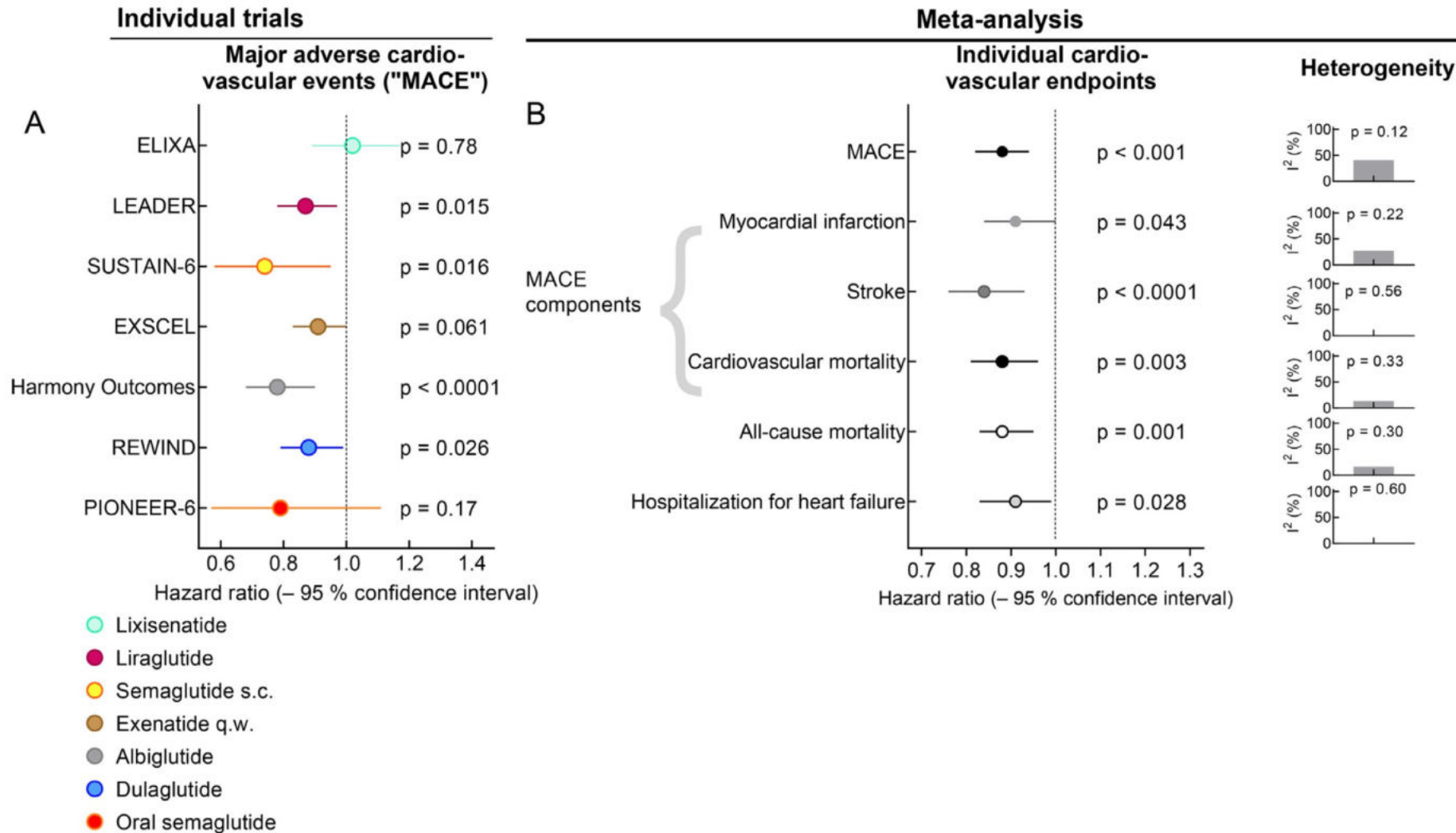
LDL Düşürücü etki



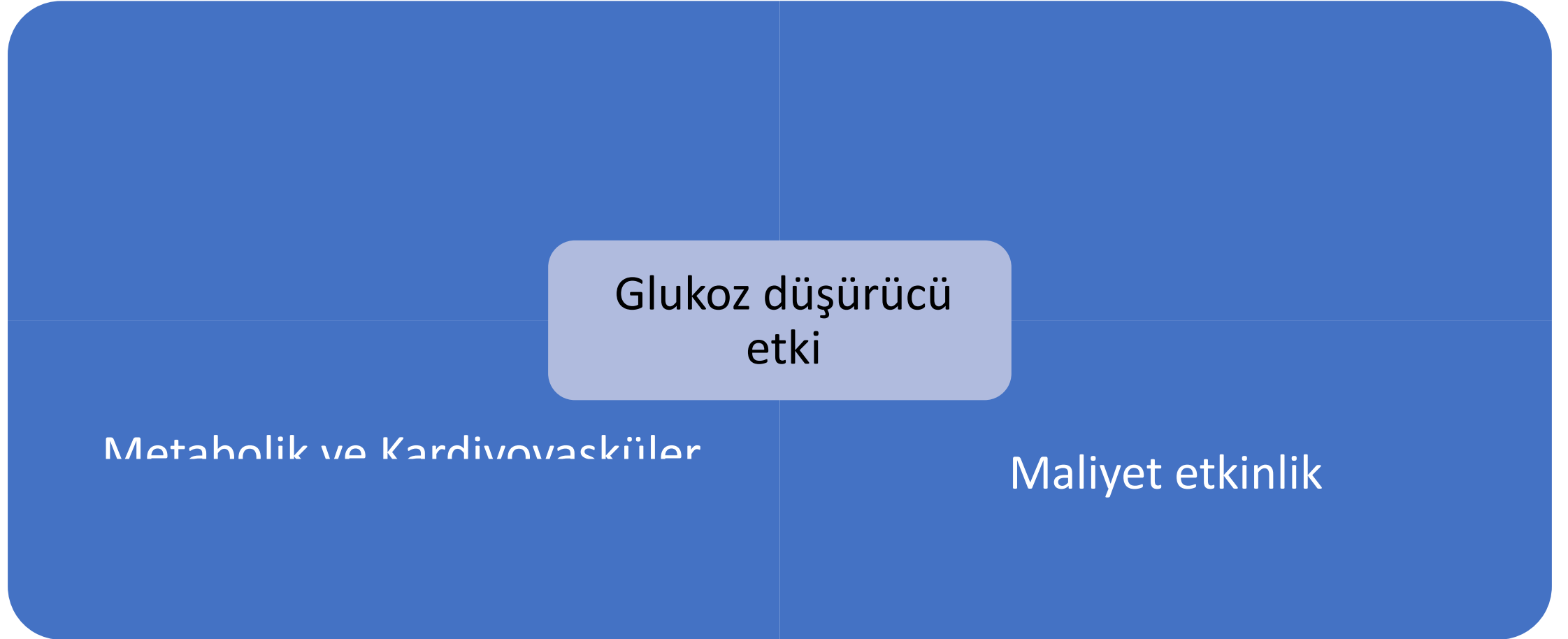
Kan basıncı üzerindeki etki



Olay ve ölümler açısından GLP-1 agonistleri



İnsülin ve GLP-1 agonisti karşılaştırma



GLP-1 agonistlerinin maliyet etkinliği insülden daha iyidir..

Open access

Original research

BMJ Open
Diabetes
Research
& Care

Glucagon-like peptide 1 agonists for treatment of patients with type 2 diabetes who fail metformin monotherapy: systematic review and meta-analysis of economic evaluation studies

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Nathorn Chaiyakunapruk,⁶ Mark McEvoy,⁷ John Attia,^{7,8} Ammarin Thakkinian^{2,9}

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ABSTRACT

Objectives To conduct a systematic review and meta-analysis and to pool the incremental net benefits (INBs) of glucagon-like peptide 1 (GLP1) compared with other therapies in type 2 diabetes mellitus (T2DM) after metformin monotherapy failure.

Research design and methods The study design is a systematic review and meta-analysis. We searched MEDLINE (via PubMed), Scopus and Tufts Registry for eligible cost-utility studies up to June 2018, adhering to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guideline. We conducted a systematic review and pooled the INBs of GLP1s compared with other

Significance of this study

What is already known about this subject?

- ▶ Glucagon-like peptide 1 (GLP1) agonists are clinically effective in treating patients with type 2 diabetes mellitus (T2DM) who fail metformin monotherapy.
- ▶ Several economic evaluation studies, along with systematic reviews of the cost-effectiveness of GLP1 agonists, have already been conducted, but these have only been descriptive and results have been conflicting.

BMJ Open Diab Res Care: first published as 10.1136/bmjdc-2019-001020 on 19 July 2020. 1



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Value of GLP-1 receptor agonists versus long-acting insulins for type 2 diabetes patients with and without established cardiovascular or chronic kidney diseases: A model-based cost-effectiveness analysis using real-world data

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GLP-1 Agonisti kullanmamak daha güvenli olabilir.

Sarkopeni

İleri karaciğer hastalığı

Pankreatit öyküsü

Yaşlı, kırılğan hasta

Akut katabolik durum

Yatan hasta

Bitirirken..

- İnsülin yaşam kurtarır, aşağıdaki durumlarda mutlaka kullanılmalıdır:
 - Tip 1 Diyabet
 - Hiperglisemik acil durumlar
 - Diyabetli bireyin perioperatif izlemi
 - Gebelik diyabeti
- Tip 2 Diyabette
 - Yaşam biçimi yönetimi ve Oral antidiyabetiklerle yeterli metabolik kontrol sağlanamazsa GLP-1 Agonisti insüline tercih edilmelidir
 - Aterosklerotik hastalık, kronik böbrek hastalığı, Obezite varsa GLP-1 agonisti (\pm Metformin) kullanılmalıdır.

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Teşekkürler..

