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Efficacy and Safety of LipiFlow® versus
Intense Pulsed Light for Meibomian
Gland Dysfunction: A Systematic Review
and Meta-analysis

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Background & Clinical Context

Meibomian gland dysfunction (MGD) : Primary cause of evaporative Dry Eye Disease (DED); marked by gland obstruction and unstable tear film.

Prevalence: Affects 30–70% globally; underdiagnosed, undertreated.

Clinical Impact: Leads to discomfort, visual disturbance, and reduced quality of life.

Treatment Gap: Traditional options (compresses, hygiene) yield temporary benefit.

Emerging Therapies: LipiFlow® and IPL (Intense Pulsed Light) are device-based treatments .

Study Rationale & Objective

Although LipiFlow® and IPL are used for MGD, direct comparisons are limited.

This study aims to

- Compare the efficacy of LipiFlow® and IPL in improving TBUT, MGS, and OSDI.
- Assess safety and patient-reported satisfaction.

Method

- . **Study design:** Systematic review & meta-analysis
- . **Study guidelines:** followed PRISMA 2020
- . **Databases:** PubMed, CENTRAL, Google Scholar (2012–2023)
- . **Eligibility criteria**
 - . **Inclusion:** English RCTs; adults with MGD; LipiFlow® vs IPL
 - . **Exclusion:** Non-RCTs, reviews, case reports, non-English

Study Selection & Outcomes

Study Selection: 13 studies; >1,200 participants; multicenter, international.

:Outcomes

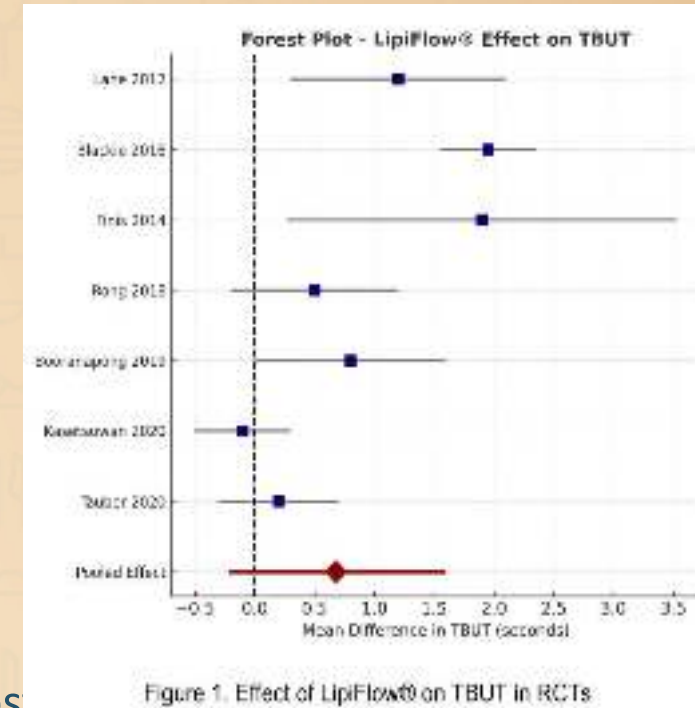
Primary: Tear Break-Up Time (TBUT), Meibomian Gland Score (MGs), Ocular Surface Disease-Index (OSDI)

Secondary: tear osmolarity, safety, patient satisfaction-

Bias Assessment: Cochrane RoB 2 tool used

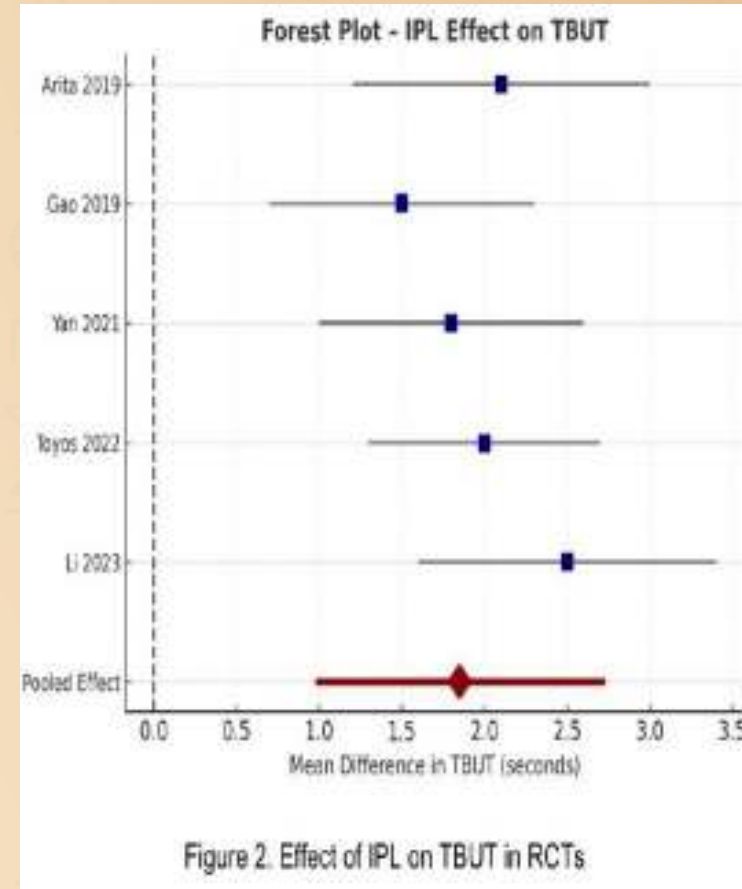
LipiFlow® : Clinical Efficacy & Safety Outcomes

- **TBUT:** Improvement up to +2.0 seconds in individual trials (e.g., Lane et al.), but pooled MD: 0.68s (95% CI: -0.22 to 1.59; $I^2 = 92.2\%$) → Not statistically significant.
- **MGS:** Strong improvement in expressibility (e.g., from 6.3 to 16.7).
- **OSDI:** Significant symptom relief (e.g., reduction from 38.1 to 24.3).
- **Tear Osmolarity:** Mild reduction in osmolarity; variable reporting across studies.
- **Patient Satisfaction:** Generally high, especially in short-term symptom improvement.
- **Safety:** highly favorable; no serious adverse events; transient pressure discomfort most common.



IPL: Clinical Efficacy & Safety Outcomes

- **TBUT:** Consistently improved across trials. Pooled MD: +1.85s (95% CI: 0.98–2.73); statistically significant.
- **MGS:** 193% increase in gland scores (Yan et al.); enhanced lipid secretion.
- **OSDI:** Sustained improvement, especially in studies with IPL + MGX (Arita et al.; Toyos et al.).
- **Tear Osmolarity:** Improved in trials using combined IPL + MGX.
- **Patient Satisfaction:** High across all time points, particularly with longer follow-up.
- **Safety:** Excellent; transient, mild erythema or warmth reported, resolving without intervention.



Comparative Analysis Table

Outcomes	LipiFlow®	IPL
Tear Break-Up Time (TBUT)	+0.68 sec (NS, pooled)	+1.85 sec (p<0.001)
Meibomian Gland Score (MGS)	↑ Significant improvement (More clear secretions)	↑ Significant improvement (better gland expressibility)
Ocular Surface Disease Index (OSDI)	↑ Symptom relief (reduced OSDI scores)	↑ Symptom relief (reduced OSDI scores)
Tear Osmolarity	Limited effect, no consistent reduction	Improved osmolarity in several trials
Patient Satisfaction	High satisfaction	High satisfaction
Safety & Tolerability	Well-tolerated, mild transient discomfort	Well-tolerated, mild transient erythema

Table 1. Comparative safety and effectiveness of LipiFlow® and IPL in MGD

Limitations & Future Directions

:Limitations

- .Substantial **heterogeneity** ($I^2 > 90\%$ in TBUT)
- .**Short follow-up** (mostly ≤ 12 weeks)-
- .Many **small-sample RCTs** (< 50 participants)-
- .Some trials lacked **blinding** or complete reporting-
- .**Geographic clustering** (Asia-dominant studies)-

:Future Research

- Long-term, multicenter head-to-head RCTs.
- Cost-effectiveness analysis.
- Stratification by MGD subtypes.

Conclusion

- .Both LipiFlow® and IPL are **safe, effective, and evidence-supported** treatments for MGD
- .Both therapies shows **substantial and sustained improvements** in MGD
- .Clinical choice should be **individualized**
- .Combination or sequential use may provide **synergistic benefits**
- .**Need for** large-scale, multicenter, standardized RCTs with longer follow-up

References



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