

Seqüenciamento (continuação...)

Felipe R. da Silva
Embrapa Recursos Genéticos e Biotecnologia



Embrapa
Biotecnologia

Novas metodologias promissoras" (2001)

- Seqüenciamento por hibridização
 - Khrapko *et al.* (1989). *FEBS Lett.* **256**: 118-122
[http://dx.doi.org/10.1016/0014-5793\(89\)81730-2](http://dx.doi.org/10.1016/0014-5793(89)81730-2)
- Seqüenciamento paralelo de assinaturas baseado em ligação e corte (MPSS)
 - Brenner *et al.* (2000). *Nature Biot.* **18**: 630 - 634
<http://dx.doi.org/10.1038/76469>
- Piroseqüenciamento
 - Ronaghi *et al.* (1996). *Anal. Biochem.* **242**: 84–89.
<http://dx.doi.org/10.1006/abio.1996.0432>

Genome Research **11**: 3–11 (2001). <http://www.genome.org/cgi/content/full/11/1/3>

Advanced Sequencing Technology Awards 2005 (NHGRI)

- Droplet-Based Digital Microfluidic Genome Sequencing
- Single-Molecule DNA Sequencing with Engineered Nanopores
- Electronic Sequencing in Nanopores
- Real-Time DNA Sequencing
- Massively Parallel Cloning and Sequencing of DNA
- Modulating Nucleotide Size in DNA for Detection by Nanopore
- Haplotype Sequencing via Single Molecule Hybridization
- Sequencing a DNA Molecule using a Synthetic Nanopore
- Real-time Multiplex Single-Molecule DNA Sequencing
- Bead-Based Polony Sequencing
- Ultra High Throughput DNA Sequencing System Based on Two-Dimensional Monolith Multi-Capillary Arrays and Nanoliter Reaction Volume
- \$100,000 Genome Using Integrated Microfluidic CE

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tecnologias de seqüenciamento de "próxima" geração

- 454 (Roche)
- SOLiD (Applied Biosystems)
- Solexa (Illumina)

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Genome Sequencer 20 System

Roche Applied Science
454 Life Sciences

Genome sequencing in microfabricated high-density picolitre reactors.
Nature **437**, 376-380
(15 September 2005)

The apparatus (...) is able to sequence 25 million bases, at 99% or better accuracy, in one 4-hour run. (...) shotgun sequencing and *de novo* assembly of the *Mycoplasma genitalium* [580 Mb] genome with 96% coverage at 99.96% accuracy in one run of the machine.

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<http://dx.doi.org/10.1038/nature03959>

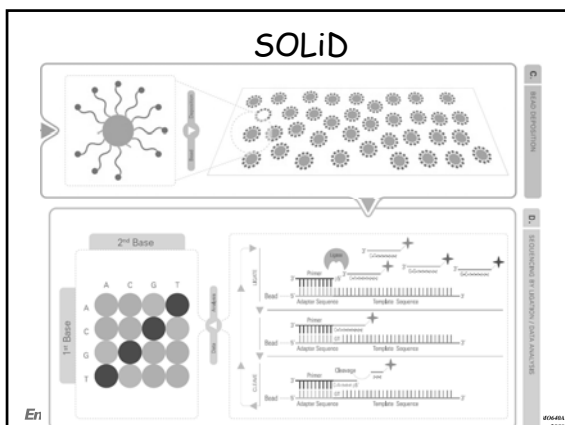
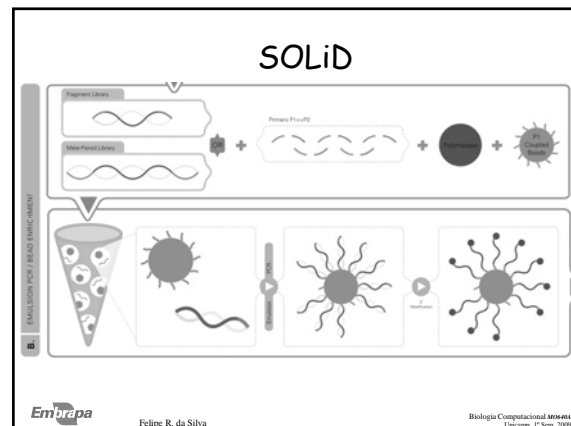
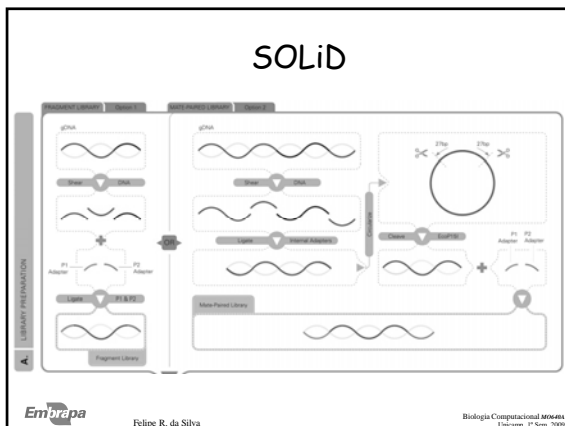
Piroseqüenciamento

$$\text{DNA}_n + \text{Nucleotéido} \xrightarrow{\text{Polimerase}} \text{DNA}_{n+1} + \text{PPi}$$

$$\text{PPi} + \text{APS} \xrightarrow{\text{ATP Sulforilase}} \text{SO}_4 + \text{ATP}$$

$$\text{ATP} + \text{Luciferina} + \text{O}_2 \xrightarrow{\text{Luciferase}} \text{AMP} + \text{PPi} + \text{Oxiluciferina} + \text{CO}_2 + \text{Luz}$$

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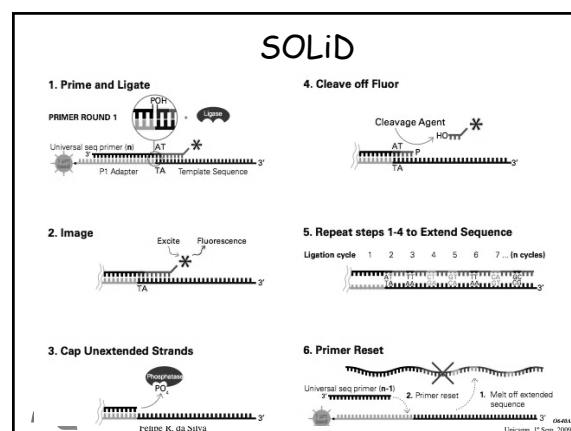
Oligo

- 2 primeiros nucleotídeos
 - dinucleotídeo específico
- nucleotídeos 3-5
 - degenerados
- nucleotídeos 6-8 *pode ser inosina.*

CGNNNI I I

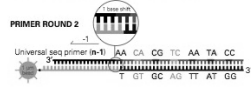
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- ### Procedimento
1. Anela *primer* e liga oligo
 2. Trata as extremidades
 3. Captura a fluorescência
 4. Remove o fluoróforo
 - Deixa um pentanucleotídeo ligado
 5. Repete o procedimento (5-7x)
 6. Denatura
 7. Utiliza um novo primer
 - (comprimento n-1)
- Embrapa* Felipe R. da Silva Biologia Computacional M06464 Volume 1, 2 Sem. 2022

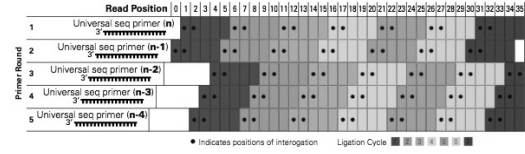


SOLiD

7. Repeat steps 1-5 with new primer

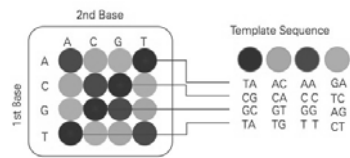


8. Repeat Reset with , n-2, n-3, n-4 primers



SOLiD

Possible Dinucleotides Encoded By Each Color



Double Interrogation



SOLiD

